

```

Matches 55; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 GCGGTGACGGTGGAGGCTATATAGCAGAGCTCGTTTAGTGAACCGTCACACCGTC 60
|||||
Db 228 GCGGTGACGGTGGAGGCTATATAGCAGAGCTCGTTTAGTGAACCGTCAGATCGCC 170

RESULT 7
BQ248646/c
LOCUS BQ248646
DEFINITION BQ248646 566 bp mRNA linear EST 03-MAY-2002
TAE25006C10F Tae25 Triticum aestivum cDNA clone Tae25006C10F, mRNA
sequence.
ACCESSION BQ248646
VERSION BQ248646.1 GI:20444522
KEYWORDS EST.
SOURCE bread wheat.
ORGANISM Triticum aestivum
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae
; Triticeae; Triticum.
1 (bases 1 to 566)
Cloutier, S.
Wheat functional genomics - Glenlea developing seeds cDNA libraries
Unpublished (2002)
Contact: Dr. Sylvie Cloutier
Cereal Research Centre, Agriculture and Agri-food Canada
195 Dafoe Rd, Winnipeg, MB, Canada R3T 2M9
Tel: (204) 983-2340
Fax: (204) 983-4604
Email: scloutier@em.agr.ca
was cloned directionally, not all sequences generated with reverse
primer were from the 5' end (same with forward primer and 3' end).
Average insert size is >870 bp
Plate: 006 row: C column: 10
Seq primer: M13 Forward.
FEATURES
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Location/Qualifiers
/organism="Triticum aestivum"
/cultivar="Glenlea"
/db_xref="taxon:4565"
/clone="Tae25006C10F"
/tissue_type="developing seeds"
/dev_stage="25 days after anthesis"
/lab_host="E. coli DH10B"
/notes="Vector: pCMV-SPORT6.0 (Invitrogen Technologies);
Site_1: NotI; Site_2: MluI; mRNA obtained from wheat seeds
of cultivar Glenlea 25 days post-anthesis"
BASE COUNT 136 a 138 c 143 g 149 t
ORIGIN
1..566
Query Match 71.1%; Score 52.6; DB 14; Length 566;
Best Local Similarity 93.2%; Pred. No. 5.2e-09;
Matches 55; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 GCGGTGACGGTGGAGGCTATATAGCAGAGCTCGTTTAGTGAACCGTCACACCGTC 60
|||||
Db 228 GCGGTGACGGTGGAGGCTATATAGCAGAGCTCGTTTAGTGAACCGTCAGATCGCC 170

RESULT 8
BQ248555/c
LOCUS BQ248555
DEFINITION BQ248555 575 bp mRNA linear EST 03-MAY-2002
Tae25007E01F Tae25 Triticum aestivum cDNA clone Tae25007E01F, mRNA
sequence.
ACCESSION BQ248555
VERSION BQ248555.1 GI:20444431
KEYWORDS EST.
SOURCE bread wheat.
ORGANISM Triticum aestivum
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae
; Triticeae; Triticum.
1 (bases 1 to 575)
Cloutier, S.
Wheat functional genomics - Glenlea developing seeds cDNA libraries
Unpublished (2002)
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Tel: (204) 983-2340
Fax: (204) 983-4604
Email: scloutier@em.agr.ca
was cloned directionally, not all sequences generated with reverse
primer were from the 5' end (same with forward primer and 3' end).
Average insert size is >870 bp
Plate: 013 row: F column: 06
Seq primer: M13 Forward.
FEATURES
source
Location/Qualifiers
/organism="Triticum aestivum"
/cultivar="Glenlea"
/db_xref="taxon:4565"
/clone="Tae25007E01F"
/tissue_type="developing seeds"
/dev_stage="25 days after anthesis"
/lab_host="E. coli DH10B"
/notes="Vector: pCMV-SPORT6.0 (Invitrogen Technologies);
Site_1: NotI; Site_2: MluI; mRNA obtained from wheat seeds
of cultivar Glenlea 25 days post-anthesis"
BASE COUNT 136 a 138 c 143 g 149 t
ORIGIN
1..566
Query Match 71.1%; Score 52.6; DB 14; Length 566;
Best Local Similarity 93.2%; Pred. No. 5.2e-09;
Matches 55; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 GCGGTGACGGTGGAGGCTATATAGCAGAGCTCGTTTAGTGAACCGTCACACCGTC 60
|||||
Db 228 GCGGTGACGGTGGAGGCTATATAGCAGAGCTCGTTTAGTGAACCGTCAGATCGCC 170

RESULT 9
BQ248056/c
LOCUS BQ248056
DEFINITION BQ248056 593 bp mRNA linear EST 03-MAY-2002
Tae25013F06F Tae25 Triticum aestivum cDNA clone Tae25013F06F, mRNA
sequence.
ACCESSION BQ248056
VERSION BQ248056.1 GI:20443932
KEYWORDS EST.
SOURCE bread wheat.
ORGANISM Triticum aestivum
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae
; Triticeae; Triticum.
1 (bases 1 to 593)
Cloutier, S.
Wheat functional genomics - Glenlea developing seeds cDNA libraries
Unpublished (2002)
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Tel: (204) 983-2340
Fax: (204) 983-4604
Email: scloutier@em.agr.ca
was cloned directionally, not all sequences generated with reverse
primer were from the 5' end (same with forward primer and 3' end).
Average insert size is >870 bp
Plate: 013 row: F column: 06
Seq primer: M13 Forward.
FEATURES
source
Location/Qualifiers
/organism="Triticum aestivum"
/cultivar="Glenlea"
/db_xref="taxon:4565"
/clone="Tae25013F06F"
/tissue_type="developing seeds"
/dev_stage="25 days after anthesis"
/lab_host="E. coli DH10B"
/notes="Vector: pCMV-SPORT6.0 (Invitrogen Technologies);
Site_1: NotI; Site_2: MluI; mRNA obtained from wheat seeds
of cultivar Glenlea 25 days post-anthesis"
BASE COUNT 141 a 140 c 142 g 151 t
ORIGIN
1..575
Query Match 71.1%; Score 52.6; DB 14; Length 575;
Best Local Similarity 93.2%; Pred. No. 5.3e-09;
Matches 55; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 GCGGTGACGGTGGAGGCTATATAGCAGAGCTCGTTTAGTGAACCGTCACACCGTC 60
|||||
Db 230 GCGGTGACGGTGGAGGCTATATAGCAGAGCTCGTTTAGTGAACCGTCAGATCGCC 172

RESULT 9
BQ248056/c
LOCUS BQ248056
DEFINITION BQ248056 593 bp mRNA linear EST 03-MAY-2002
Tae25013F06F Tae25 Triticum aestivum cDNA clone Tae25013F06F, mRNA
sequence.
ACCESSION BQ248056
VERSION BQ248056.1 GI:20443932
KEYWORDS EST.
SOURCE bread wheat.
ORGANISM Triticum aestivum
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae
; Triticeae; Triticum.
1 (bases 1 to 593)
Cloutier, S.
Wheat functional genomics - Glenlea developing seeds cDNA libraries
Unpublished (2002)
Contact: Dr. Sylvie Cloutier
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195 Dafoe Rd, Winnipeg, MB, Canada R3T 2M9
Tel: (204) 983-2340
Fax: (204) 983-4604
Email: scloutier@em.agr.ca
was cloned directionally, not all sequences generated with reverse
primer were from the 5' end (same with forward primer and 3' end).
Average insert size is >870 bp
Plate: 013 row: F column: 06
Seq primer: M13 Forward.
FEATURES
source
Location/Qualifiers
/organism="Triticum aestivum"
/cultivar="Glenlea"
/db_xref="taxon:4565"
/clone="Tae25013F06F"
/tissue_type="developing seeds"
/dev_stage="25 days after anthesis"
/lab_host="E. coli DH10B"
/notes="Vector: pCMV-SPORT6.0 (Invitrogen Technologies);
Site_1: NotI; Site_2: MluI; mRNA obtained from wheat seeds
of cultivar Glenlea 25 days post-anthesis"
BASE COUNT 141 a 140 c 142 g 151 t
ORIGIN
1..575
Query Match 71.1%; Score 52.6; DB 14; Length 575;
Best Local Similarity 93.2%; Pred. No. 5.3e-09;
Matches 55; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 GCGGTGACGGTGGAGGCTATATAGCAGAGCTCGTTTAGTGAACCGTCACACCGTC 60
|||||
Db 230 GCGGTGACGGTGGAGGCTATATAGCAGAGCTCGTTTAGTGAACCGTCAGATCGCC 172

```

```

/clone="Tae25013F06F"
/clone_lib="Tae25"
/tissue_type="developing seeds"
/dev_stage="25 days after anthesis"
/lab_host="E. coli DH10B"
/notes="vector: pCMV-SPORT6.0 (Invitrogen Technologies);
Site_1: NotI; Site_2: MluI; mRNA obtained from wheat seeds
of cultivar Glenlea 25 days post-anthesis"

BASE COUNT      145 a   146 c   147 g   155 t
ORIGIN
Query Match      71.1%; Score 52.6; DB 14; Length 593;
Best Local Similarity 93.2%; Pred. No. 5.3e-09;
Matches 55; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Oy 2 GCGGTGTCGGTGGGAGGCTATATAAGCAGAGCTGTTTACTGTGAACCGTCAACCGTC 60
|||||
b 228 GCGGTGTCGGTGGGAGGCTATATAAGCAGAGCTGTTTACTGTGAACCGTCAACCGTC 170
|||||

RESULT 10
B68191/c
LOCUS
DEFINITION      B68191 610 bp DNA linear GSS 18-JUN-1998
CIT978SK-A-492C12.TV CIT978SK Homo sapiens genomic clone A-492C12,
DNA sequence.
ACCESSION      B68191
VERSION        B68191.1 GI:2666901
KEYWORDS      GSS.
SOURCE        human.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 610)
Adams M.D., Rounsley S.D., Field C.E., Bass S., Linher K., Golden
K., Berry K., Granger D., Suh E., Wible C., Kim U.-J., Shizuya H.,
Simon M. and Venter J.C.
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: mdadams@tigr.org
Clones are available from Research Genetics (info@resgen.com). BAC
end search page:
http://www.tigr.org/tldb/hungen/bac_end_search/bac_end_search.html
Seq primer: 77
Class: BAC ends.

FEATURES
Location/Qualifiers
1..610
/organism="Homo sapiens"
/db_xref="GDB:726476"
/db_xref="taxon:9606"
/clone="A-492C12"
/clone_lib="CIT978SK"
/sex="Female"
/cell_type="Fibroblast"
/notes="Vector: pBAC108L; Site_1: HindIII; Site_2: HindIII;
Caltech Human BAC Library A"

BASE COUNT      146 a   159 c   165 g   140 t
ORIGIN
Query Match      71.1%; Score 52.6; DB 17; Length 610;
Best Local Similarity 93.2%; Pred. No. 5.4e-09;
Matches 55; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Oy 2 GCGGTGTCGGTGGGAGGCTATATAAGCAGAGCTGTTTACTGTGAACCGTCAACCGTC 60
|||||
b 288 GCGGTGTCGGTGGGAGGCTATATAAGCAGAGCTGTTTACTGTGAACCGTCAACCGTC 230
|||||

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RESULT 11
BQ248255/c
LOCUS
DEFINITION      BQ248255 625 bp mRNA linear EST 03-MAY-2002
Tae25011C08F Tae25 Triticum aestivum cDNA clone Tae25011C08F, mRNA
sequence.
ACCESSION      BQ248255
VERSION        BQ248255.1 GI:20444131
KEYWORDS      EST.
SOURCE        bread wheat.
ORGANISM      Triticum aestivum
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae
; Triticeae; Triticum.
1 (bases 1 to 625)
Cloutier, S.
Wheat functional genomics - Glenlea developing seeds cDNA libraries
Unpublished (2002)
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195 Dafoe Rd, Winnipeg, MB, Canada R3T 2M9
Tel: (204) 983-2340
Fax: (204) 983-4604
Email: scloutier@agr.ca
was cloned directionally, not all sequences generated with reverse
primer were from the 5' end (same with forward primer and 3' end).
Average insert size is >870 bp
Plate: 011 row: C column: 08
Seq primer: M13 Forward.

FEATURES
Location/Qualifiers
1..625
/organism="Triticum aestivum"
/cultivar="Glenlea"
/db_xref="taxon:4565"
/clone="Tae25011C08F"
/clone_lib="Tae25"
/tissue_type="developing seeds"
/dev_stage="25 days after anthesis"
/lab_host="E. coli DH10B"
/notes="Vector: pCMV-SPORT6.0 (Invitrogen Technologies);
Site_1: NotI; Site_2: MluI; mRNA obtained from wheat seeds
of cultivar Glenlea 25 days post-anthesis"

BASE COUNT      155 a   152 c   154 g   164 t
ORIGIN
Query Match      71.1%; Score 52.6; DB 14; Length 625;
Best Local Similarity 93.2%; Pred. No. 5.4e-09;
Matches 55; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Oy 2 GCGGTGTCGGTGGGAGGCTATATAAGCAGAGCTGTTTACTGTGAACCGTCAACCGTC 60
|||||
b 228 GCGGTGTCGGTGGGAGGCTATATAAGCAGAGCTGTTTACTGTGAACCGTCAACCGTC 170
|||||

RESULT 12
BQ248467/c
LOCUS
DEFINITION      BQ248467 625 bp mRNA linear EST 03-MAY-2002
Tae25008F09F Tae25 Triticum aestivum cDNA clone Tae25008F09F, mRNA
sequence.
ACCESSION      BQ248467
VERSION        BQ248467.1 GI:20444343
KEYWORDS      EST.
SOURCE        bread wheat.
ORGANISM      Triticum aestivum
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae
; Triticeae; Triticum.
1 (bases 1 to 625)
Cloutier, S.
Wheat functional genomics - Glenlea developing seeds cDNA libraries
Unpublished (2002)
Contact: Dr. Sylvie Cloutier

```

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195 Dafoe Rd, Winnipeg, MB, Canada R3T 2M9  
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Fax: (204) 983-4604  
Email: scloutier@em.agr.ca

was cloned directionally, not all sequences generated with reverse primer were from the 5' end (same with forward primer and 3' end).  
Average insert size is >870 bp  
Plate: 008 row: F column: 09  
Seq primer: M13 Forward.

## FEATURES

## source

1. .625  
/organism="Triticum aestivum"  
/cultivar="Glenlea"  
/db\_xref="taxon:4565"  
/clone="TaE25008F09F"  
/clone\_lib="TaE25"  
/tissue\_type="developing seeds"  
/dev\_stage="25 days after anthesis"  
/lab\_host="E. coli DH10B"  
/note="Vector: pCMV-SPORT6.0 (Invitrogen Technologies); Site\_1: NotI; Site\_2: MluI; mRNA obtained from wheat seeds of cultivar Glenlea 25 days post-anthesis"

E COUNT

155 a 152 c 155 g 163 t

## ..IGIN

Query Match 71.1%; Score 52.6; DB 14; Length 625;  
Best Local Similarity 93.2%; Pred. No. 5.4e-09;  
Matches 55; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 GCGGTGACGGTGGGAGGCTATATAAGCAGAGCTCGTTTACTGAACCGTCAACCGTC 60  
|||||  
Db 228 GCGGTGACGGTGGGAGGCTATATAAGCAGAGCTCGTTTACTGAACCGTCAACCGTC 170

## RESULT 13

BQ248453/c

LOCUS

DEFINITION BQ248453 630 bp mRNA linear EST 03-MAY-2002  
TaE25008H04F TaE25 Triticum aestivum cDNA clone TaE25008H04F, mRNA sequence.

ACCESSION

BQ248453

VERSION

BQ248453.1

KEYWORDS

EST.

SOURCE

bread wheat.

ORGANISM

Triticum aestivum

AUTHORS

Cloutier,S.

TITLE

Unpublished (2002)

JOURNAL

COMMENT

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Tel: (204) 983-2340

Fax: (204) 983-4604

Email: scloutier@em.agr.ca

was cloned directionally, not all sequences generated with reverse primer were from the 5' end (same with forward primer and 3' end).

Average insert size is &gt;870 bp

Plate: 008 row: H column: 04

Seq primer: M13 Forward.

Location/Qualifiers

1. .630

/organism="Triticum aestivum"

/cultivar="Glenlea"

/db\_xref="taxon:4565"

/clone="TaE25008H04F"

/clone\_lib="TaE25"

/tissue\_type="developing seeds"

/dev\_stage="25 days after anthesis"

/lab\_host="E. coli DH10B"

## FEATURES

## source

1

/note="Vector: pCMV-SPORT6.0 (Invitrogen Technologies); Site\_1: NotI; Site\_2: MluI; mRNA obtained from wheat seeds of cultivar Glenlea 25 days post-anthesis"  
BASE COUNT 156 a 154 c 155 g 165 t  
ORIGIN

Query Match 71.1%; Score 52.6; DB 14; Length 630;  
Best Local Similarity 93.2%; Pred. No. 5.5e-09;  
Matches 55; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
QY 2 GCGGTGACGGTGGGAGGCTATATAAGCAGAGCTCGTTTACTGAACCGTCAACCGTC 60  
|||||  
Db 228 GCGGTGACGGTGGGAGGCTATATAAGCAGAGCTCGTTTACTGAACCGTCAACCGTC 170

## RESULT 14

BQ248173/c

LOCUS

DEFINITION BQ248173 631 bp mRNA linear EST 03-MAY-2002  
TaE25012C11F TaE25 Triticum aestivum cDNA clone TaE25012C11F, mRNA sequence.

ACCESSION

BQ248173

VERSION

BQ248173.1

KEYWORDS

EST.

SOURCE

bread wheat.

ORGANISM

Triticum aestivum

AUTHORS

Cloutier,S.

TITLE

Unpublished (2002)

JOURNAL

COMMENT

Contact: Dr. Sylvie Cloutier

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Fax: (204) 983-4604

Email: scloutier@em.agr.ca

was cloned directionally, not all sequences generated with reverse primer were from the 5' end (same with forward primer and 3' end).

Average insert size is &gt;870 bp

Plate: 012 row: C column: 11

Seq primer: M13 Forward.

Location/Qualifiers

1. .631

/organism="Triticum aestivum"

/cultivar="Glenlea"

/db\_xref="taxon:4565"

/clone="TaE25012C11F"

/clone\_lib="TaE25"

/tissue\_type="developing seeds"

/dev\_stage="25 days after anthesis"

/lab\_host="E. coli DH10B"

/note="Vector: pCMV-SPORT6.0 (Invitrogen Technologies); Site\_1: NotI; Site\_2: MluI; mRNA obtained from wheat seeds of cultivar Glenlea 25 days post-anthesis"

BASE COUNT 156 a 154 c 155 g 166 t

ORIGIN

## Query Match

Best Local Similarity 93.2%; Pred. No. 5.5e-09;  
Matches 55; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 GCGGTGACGGTGGGAGGCTATATAAGCAGAGCTCGTTTACTGAACCGTCAACCGTC 60  
|||||  
Db 228 GCGGTGACGGTGGGAGGCTATATAAGCAGAGCTCGTTTACTGAACCGTCAACCGTC 170

## RESULT 15

BQ248939/c

LOCUS

DEFINITION BQ248939 633 bp mRNA linear EST 03-MAY-2002  
TaE25002C05F TaE25 Triticum aestivum cDNA clone TaE25002C05F, mRNA sequence.

ACCESSION BQ248939  
 VERSION BQ248939.1 GI:20444815  
 KEYWORDS EST.  
 SOURCE bread wheat.  
 ORGANISM *Triticum aestivum*  
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae  
 ; Triticeae; *Triticum*.  
 1 (bases 1 to 633)  
 Cloutier, S.  
 Wheat functional genomics - Glenlea developing seeds cDNA libraries  
 Unpublished (2002)  
 Contact: Dr. Sylvie Cloutier  
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 Tel: (204) 983-2340  
 Fax: (204) 983-4604  
 Email: scloutier@em.agr.ca  
 was cloned directionally, not all sequences generated with reverse  
 primer were from the 5' end (same with forward primer and 3' end).  
 Average insert size is >870 bp  
 Plate: 002 row: C column: 05  
 Seq primer: M13 Forward.

# FEATURES source

1..633  
 /organism="Triticum aestivum"  
 /cultivar="Glenlea"  
 /db\_xref="taxon:4565"  
 /clone="TaE25002C05F"  
 /clone\_lib="TaE25"  
 /tissue\_type="developing seeds"  
 /dev\_stage="25 days after anthesis"  
 /lab\_host="E. coli DH10B"  
 /note="Vector: pCMV-SPORT6.0 (Invitrogen Technologies);  
 Site\_1: NotI; Site\_2: MluI; mRNA obtained from wheat seeds  
 of cultivar Glenlea 25 days post-anthesis"  
 157 a 154 c 154 g 168 t

# BASE COUNT ORIGIN

Query Match 71.1%; Score 52.6; DB 14; Length 633;  
 Best Local Similarity 93.2%; Pred. NO. 5.5e-09;  
 Matches 55; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
 Qy 2 GCGGTGTACGGTGGGAGGCTATATAGCAGAGCTCGTTAGTGAACCGTCAACCGTC 60  
 ||||||||||||||| ||||||||||||||||||||||||||||||| |||  
 b 228 GCGGTGTACGGTGGGAGGCTATATAGCAGAGCTCGTTAGTGAACCGTCAACCGTC 170

Search completed: March 6, 2003, 00:22:23  
 Job time : 560.372 secs



Plate: 014 row: A column: 07  
Seq primer: M13 Forward.  
Location/Qualifiers  
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/organism="Triticum aestivum"  
/cultivar="Glenlea"  
/db\_xref="taxon:4565"  
/clone="TaE25014A07F"  
/clone\_lib="TaE25"  
/tissue\_type="developing seeds"  
/dev\_stage="25 days after anthesis"  
/lab\_host="E. coli DH10B"  
/note="Vector: pCMV-SPORT6.0 (Invitrogen Technologies);  
Site\_1: NotI; Site\_2: MluI; mRNA obtained from wheat seeds  
of cultivar Glenlea 25 days post-anthesis"

BASE COUNT 121 a 128 c 122 g 131 t  
ORIGIN

Query Match 71.1%; Score 52.6; DB 14; Length 502;  
Best Local Similarity 93.2%; Pred. No. 5e-09;  
Matches 55; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 2 GCGGTGACGGTGGAGGCGCTATATAGCAGAGCTCGTTTAGTGAACCGTCAAAACCGTC 60  
|||||  
Db 228 GCGGTGACGGTGGAGGCGCTATATAGCAGAGCTCGTTTAGTGAACCGTCAAGTCGCC 170  
|||||

RESULT 6  
LOCUS BQ248278/c  
DEFINITION TaE25011A03F TaE25 Triticum aestivum cDNA clone TaE25011A03F, mRNA  
sequence.  
ACCESSION BQ248278  
VERSION BQ248278.1 GI:20444154  
KEYWORDS EST.  
SOURCE bread wheat.  
ORGANISM Triticum aestivum  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae;  
: Triticeae; Triticum.  
1 (bases 1 to 535)  
Cloutier,S.  
Wheat functional genomics - Glenlea developing seeds cDNA libraries  
Unpublished (2002)  
Contact: Dr. Sylvie Cloutier  
Cereal Research Centre, Agriculture and Agri-food Canada  
195 Daffoe Rd, Winnipeg, MB, Canada R3T 2M9  
Tel: (204) 983-2340  
Fax: (204) 983-4604  
Email: scloutier@em.agr.ca  
was cloned directionally, not all sequences generated with reverse  
primer were from the 5' end (same with forward primer and 3' end).  
Average insert size is >870 bp  
Plate: 011 row: A column: 03  
Seq primer: M13 Forward.  
Location/Qualifiers  
1. .535  
/organism="Triticum aestivum"  
/cultivar="Glenlea"  
/db\_xref="taxon:4565"  
/clone="TaE25011A03F"  
/clone\_lib="TaE25"  
/tissue\_type="developing seeds"  
/dev\_stage="25 days after anthesis"  
/lab\_host="E. coli DH10B"  
/note="Vector: pCMV-SPORT6.0 (Invitrogen Technologies);  
Site\_1: NotI; Site\_2: MluI; mRNA obtained from wheat seeds  
of cultivar Glenlea 25 days post-anthesis"

BASE COUNT 130 a 133 c 132 g 139 t 1 others  
ORIGIN

Query Match 71.1%; Score 52.6; DB 14; Length 535;  
Best Local Similarity 93.2%; Pred. No. 5.1e-09;

```

FEATURES
  source
    Seq primer: T7.
    Location/Qualifiers
      1..657
        /organism="Danio rerio"
        /db_xref="taxon:7955"
        /clone="R2PD clone CHBop576F21214Q3"
        /clone_lib="zebrafish kidney cDNA random primed, R2PD
        library no: 576"
        /dev_stage="adult"
        /note="Organ: kidney; Vector: pBK-CMV"
BASE COUNT      163 a   165 c   162 g   165 t       2 others
ORIGIN
  Query Match
    71.9%; Score 53.2; DB 9; Length 657;
  Best Local Similarity
    87.9%; Pred. No. 3.3e-09;
  Matches 58; Conservative 0; Mismatches 8; Indels 0; Gaps 0;

QY 2 GGCCTGTACGGTGGAGGCTATATAGCAGAGCTCGTTTGTAGTGAACCGTCAAAACCGTCA 61
      |||||
Db 369 GGCCTGTACGGTGGAGGCTCTATATAGCAGAGCTCGTTTGTAGTGAACCGTCAAGATCCGCT 310
      |||||

QY 62 AACCGC 67
      |||||
Db 309 AGCGCG 304

RESULT 2
BE661741/c
LOCUS
DEFINITION
  138t7 GmaxSC Glycine max cDNA, mRNA sequence.
ACCESSION
  BE661741
VERSION
  BE661741.1 GI:9987633
KEYWORDS
  EST.
SOURCE
  soybean.
  ORGANISM
    Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
    Spermatophyta; Magnoliophyta; eudicotyledons; Core eudicots;
    Rosidae; eurosids I; Fabales; Fabaceae; Papilionoideae; Phaseoleae;
    Glycine.
  1 (bases 1 to 703)
  Harris, N., Chapman, B.P. and Gijzen, M.
  Gene expression in developing soybean seed coats
  Unpublished (2000)
  Contact: Gijzen M
  Agriculture and Agri-Food Canada
  1391 Sandford Street, London, Ontario, Canada N5V 4T3
  Tel: 519 457 1470
  Fax: 519 457 3997
  Email: gijzenm@agr.ca.
  Location/Qualifiers
    1..703
      /organism="Glycine max"
      /cultivar="Harosoy 63"
      /db_xref="taxon:3847"
      /clone_lib="GmaxSC"
      /tissue_type="Seed coats"
      /lab_host="E. coli strain XL0LR"
      /note="Vector: pBK-CMV; Site_1: EcoRI; Site_2: XhoI; This
      cDNA library was constructed from polyA+ enriched mRNA
      from green seed coats in mid to late developmental stage
      , average fresh weight 250 mg per seed. Traces of pod and
      embryo tissue also present. Complementary DNA was
      synthesized from mRNA using an XhoI-poly(dT)
      linker-primer. EcoRI adapters were ligated to the
      blunt-ended cDNA fragments and the products were digested
      with XhoI for directional cloning into lambda ZAP Express
      vector. This lambda library was amplified once using E.
      coli host strain XL1 Blue MRF+. Inserts were then
      subcloned by mass excision using ExAssist helper phage for
      conversion into phagemid vector pBK-CMV in E. coli host
      strain XL0LR."
BASE COUNT      199 a   147 c   148 g   203 t       6 others
ORIGIN

```

```

Query Match
  71.9%; Score 53.2; DB 10; Length 703;
  Best Local Similarity
    87.9%; Pred. No. 3.4e-09;
  Matches 58; Conservative 0; Mismatches 8; Indels 0; Gaps 0;

QY 2 GCGGTGTACGGTGGAGGCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAAAACCGTCA 61
      |||||
Db 535 GCGGTGTACGGTGGAGGCTCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAAGATCCGCT 476
      |||||

QY 62 AACCGC 67
      |||||
Db 475 AGCGCG 470

```

```

RESULT 3
B51582/c
LOCUS
DEFINITION
  C19797SK-A-481E4.TV C19797SK Homo sapiens genomic clone A-481E4,
  DNA sequence.
ACCESSION
  B51582
VERSION
  B51582.1 GI:2603819
KEYWORDS
  GSS.
SOURCE
  human.
  ORGANISM
    Homo sapiens

```

```

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
  1 (bases 1 to 329)
  Kim, U.-J., Adams, M.D. and Simon, M.I.
  Determination of clone end sequences of human Bacterial Artificial
  Chromosomes
  Unpublished (1997)
  Other_GSSs: C19797SK-A-481E4.TP
  Contact: Ung-Jin Kim
  Caltech Genome Research Lab
  California Institute of Technology
  Division of Biology, MS 147-75, Pasadena, CA 91125, USA
  Tel: 626 796 7066
  Fax: 626 395 4901
  Email: ung@ash.tree.caltech.edu
  Clones are available from Research Genetics (info@resgen.com). BAC
  end search page:
  http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html
  Seq primer: T7
  Class: BAC ends.
  Location/Qualifiers
    1..329
      /organism="Homo sapiens"
      /db_xref="GDB:726891"
      /db_xref="taxon:9606"
      /clone="A-481E4"
      /clone_lib="C19797SK"
      /sex="Female"
      /cell_type="Fibroblast"
      /note="Vector: pBAC108L; Site_1: HindIII; Site_2: HindIII;
      CalTech Human BAC Library A"
BASE COUNT      72 a   89 c   90 g   78 t
ORIGIN

```

```

Query Match
  71.1%; Score 52.6; DB 17; Length 329;
  Best Local Similarity
    93.2%; Pred. No. 4.2e-09;
  Matches 55; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 GCGGTGTACGGTGGAGGCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAAAACCGTC 60
      |||||
Db 242 GCGGTGTACGGTGGAGGCTCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAAGATCCGCC 184
      |||||

RESULT 4
BQ248969/c
LOCUS
DEFINITION
  TAE25001G10F TAE25 Triticum aestivum cDNA clone TAE25001G10F, mRNA
  sequence.
ACCESSION
  BQ248969

```

GenCore version 5.1.3  
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: March 5, 2003, 22:43:11 ; Search time 558.039 seconds  
(without alignments)  
2147.640 Million cell updates/sec

Title: US-09-980-548-3

Perfect score: 74

Sequence: 1 tggcgtgacgtggaggc.....accgtcaaccgcgggaagct 74

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:\*

1: em\_estba:\*

2: em\_esthum:\*

3: em\_estin:\*

4: em\_estmu:\*

5: em\_estov:\*

6: em\_estpl:\*

7: em\_estro:\*

8: em\_htc:\*

9: gb\_esti:\*

10: gb\_est2:\*

11: gb\_htc:\*

12: gb\_est3:\*

13: gb\_est4:\*

14: gb\_est5:\*

15: em\_estfun:\*

16: em\_estom:\*

17: gb\_gss:\*

18: em\_gss\_hum:\*

19: em\_gss\_inv:\*

20: em\_gss\_pln:\*

21: em\_gss\_vrt:\*

22: em\_gss\_fun:\*

23: em\_gss\_mam:\*

24: em\_gss\_mus:\*

25: em\_gss\_other:\*

26: em\_gss\_pro:\*

27: em\_gss\_rod:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Query Match %	Score	Length	ID	Description
C 1	53.2	71.9	657	9- AI815377	AI815377 B0283860
C 2	53.2	71.9	703	10 BE661741	BE661741 138t7 Gma
C 3	52.6	71.1	329	17 B51582	B51582 CIT978SK-A-
C 4	52.6	71.1	388	14 BQ248969	BQ248969 TAE25001G
C 5	52.6	71.1	502	14 BQ248026	BQ248026 TAE25014A
C 6	52.6	71.1	535	14 BQ248278	BQ248278 TAE25011A

C 7	52.6	71.1	566	14 BQ248646	BQ248646 TAE25006C
C 8	52.6	71.1	575	14 BQ248555	BQ248555 TAE25007E
C 9	52.6	71.1	593	14 BQ248056	BQ248056 TAE25013F
C 10	52.6	71.1	610	17 B68191	B68191 CIT978SK-A-
C 11	52.6	71.1	625	14 BQ248255	BQ248255 TAE25011C
C 12	52.6	71.1	625	14 BQ248467	BQ248467 TAE25008F
C 13	52.6	71.1	630	14 BQ248453	BQ248453 TAE25008H
C 14	52.6	71.1	630	14 BQ248453	BQ248453 TAE25012C
C 15	52.6	71.1	631	14 BQ248173	BQ248173 TAE25012C
C 16	52.6	71.1	633	14 BQ248939	BQ248939 TAE25002C
C 17	52.6	71.1	633	14 BQ248504	BQ248504 TAE25008B
C 18	52.6	71.1	641	14 BQ247936	BQ247936 TAE25015B
C 19	52.6	71.1	641	14 BQ248159	BQ248159 TAE25012E
C 20	52.6	71.1	642	14 BQ247761	BQ247761 TAE25042H
C 21	52.6	71.1	644	14 BQ247949	BQ247949 TAE25015A
C 22	52.6	71.1	644	14 BQ247963	BQ247963 TAE25014G
C 23	52.6	71.1	645	14 BQ247960	BQ247960 TAE25014H
C 24	52.6	71.1	646	14 BQ247762	BQ247762 TAE25042H
C 25	52.6	71.1	646	14 BQ248341	BQ248341 TAE25010C
C 26	52.6	71.1	647	14 BQ248311	BQ248311 TAE25010F
C 27	52.6	71.1	647	14 BQ248458	BQ248458 TAE25008G
C 28	52.6	71.1	648	14 BQ247997	BQ247997 TAE25014D
C 29	52.6	71.1	648	14 BQ248418	BQ248418 TAE25009C
C 30	52.6	71.1	649	14 BQ248913	BQ248913 TAE25002F
C 31	52.6	71.1	649	14 BQ247933	BQ247933 TAE25015B
C 32	52.6	71.1	649	14 BQ248366	BQ248366 TAE25009H
C 33	51	70.5	400	17 B63118	B63118 CIT978SK-A-
C 34	51	68.9	287	17 B63084	B63084 CIT978SK-A-
C 35	51	68.9	545	14 BQ248598	BQ248598 TAE25006H
C 36	50.8	68.6	550	14 BQ248595	BQ248595 TAE25006H
C 37	50.2	67.8	607	10 AW355220	AW355220 pnf-b.pkO
C 38	50.2	67.8	617	10 AV645401	AV645401 AV645401
C 39	50.2	67.8	618	10 AV645332	AV645332 AV645332
C 40	50.2	67.8	619	10 AV645383	AV645383 AV645383
C 41	50.2	67.8	619	10 AV645377	AV645377 AV645377
C 42	50.2	67.8	619	10 AV645379	AV645379 AV645379
C 43	50.2	67.8	619	10 AV645385	AV645385 AV645385
C 44	50.2	67.8	619	10 AV645389	AV645389 AV645389
C 45	50.2	67.8	620	10 AV645369	AV645369 AV645369
					AV645380 AV645380

## ALIGNMENTS

RESULT 1  
AI815377/c  
LOCUS B0283860 zebrafish kidney cdna random primed, RZPD library no: 576  
DEFINITION Danio rerio cdna clone RZPD clone CHBOP576F21214Q3, mRNA sequence.  
ACCESSION AI815377  
VERSION AI815377.1 GI:5430923  
KEYWORDS EST.  
SOURCE zebrafish.  
ORGANISM Danio rerio

AI815377 657 bp mRNA linear EST 09-JUL-1999  
B0283860 zebrafish kidney cdna random primed, RZPD library no: 576  
Danio rerio cdna clone RZPD clone CHBOP576F21214Q3, mRNA sequence.

AI815377.1 GI:5430923  
EST.  
zebrafish.  
Danio rerio  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes  
; Cyprinidae; Danio.

1 (bases 1 to 657)  
Look,A.T. and Holloway,M.  
Zebrafish Kidney cdna  
Unpublished (1999)  
Contact: Thomas Look

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT

Dana-Farber Cancer Institute  
Pediatric Oncology Department, Mayer Building-630.; 44 Binney Street  
, Boston, MA 02115, USA  
Tel: (617) 632-5826  
Fax: (617) 632-6989  
Email: thomas.look@dfci.harvard.edu

The clone was obtained from an RZPD array made from an adult kidney  
cdna random primed library from Leonard Zon. RZPD library number  
576. This clone is available from the Resource Centre/Primary  
Database of the German Human Genome Project: RZPD (Resource  
Zentrum Primr Datenbank). <http://www.rzpd.edu>

QY 2 GCGGTGTACGGTGGAGGCGCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAAAACCGTC 60  
|||||  
Db 327 GCGGTGTACGGTGGAGGCGCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAATCGCC 385

RESULT 15  
ID AAX60047 standard; DNA: 450 BP.  
XX AAX60047;  
XX AAX60047;  
XX 04-AUG-1999 (first entry)  
XX Minimal Human cytomegalovirus promoter.  
XX Transgenic mice; transgene; tet operator-linked gene; tetracycline;  
KW mouse-active transcriptional regulatory element; mutant Tet repressor;  
KW gene therapy; genetic disease; acquired disease; cancer; viral disease;  
KW vaccination; rheumatoid arthritis; hypopituitarism; wound healing;  
KW tissue regeneration; cancer; benign prostatic hypertrophy; hemophilia;  
KW erythrocytopenia; arteriosclerosis; liver disease; Alzheimer's disease;  
KW Parkinson's disease; human disease model; ds.

Human cytomegalovirus.  
US5912411-A.  
15-JUN-1999.  
07-JUN-1995; 93US-0487472.  
07-JUN-1995; 95US-0487472.  
14-JUN-1993; 93US-0076327.  
14-JUN-1993; 93US-0076726.  
14-JUN-1994; 94US-0260452.  
01-JUL-1994; 94US-0270637.  
15-JUL-1994; 94US-0275876.  
03-FEB-1995; 95US-0383754.  
(UYHE-) UNIV HEIDELBERG.  
Bujard H, Gossen M;  
WPI; 1999-357232/30.  
Transgenic mice with inducible transgene activity useful for in vitro and in vivo protein production  
Disclosure; Column 71-72; 63pp; English.

The specification describes transgenic mice which have a transgene and a tet operator-linked gene integrated in the genome. The transgene comprises a mouse-active transcriptional regulatory element linked to a polynucleotide sequence that encodes a fusion protein which activates transcription of the tet operator-linked gene. The fusion protein comprises a mutated Tet repressor that binds a tet operator sequence in the presence of tetracycline (Tc) or a Tc analogue, linked to a polypeptide that activates transcription in eukaryotic cells. The transgenic system may be used for gene therapy to treat genes involved in genetic or acquired diseases. Gene therapy may be used to treat cancer, viral diseases, for vaccination, and to provide (Tc induced) regulated doses of a product (e.g. for the treatment or regulation of rheumatoid arthritis, hypopituitarism, wound healing and tissue regeneration, cancer, benign prostatic hypertrophy, hemophilia, erythrocytopenia, arteriosclerosis and liver disease, Alzheimer's disease, and Parkinson's disease). The system may also be used to produce proteins in vivo (e.g. using mammalian, yeast or fungal cells) or in vitro (e.g. transgenic farm animals), to produce animal models of human disease, or to produce a stable cell line for gene cloning. The present sequence is used to construct the transgenes of the invention.

Sequence 450 BP; 104 A; 109 C; 102 G; 135 T; 0 other;

Query Match 73.2%; Score 54.2; DB 20; Length 450;  
Best Local Similarity 94.9%; Pred. No. 4.3e-12;  
Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
QY 2 GCGGTGTACGGTGGAGGCGCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAAAACCGTC 60  
|||||  
Db 327 GCGGTGTACGGTGGAGGCGCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAATCGCC 385

Search completed: March 5, 2003, 22:49:57  
Job time : 73.2126 secs



PI Bujard H, Gossen M;  
 XX WPI; 1998-541795/46.  
 XX  
 PT Tetracycline based regulation of gene expression - uses a  
 PT tetracycline operator sequence joined to a gene of interest, the  
 PT gene of interest being induced in the presence, but not absence of  
 PT the antibiotic  
 XX  
 PS Example 6; Columns 71-72; 63pp; English.  
 XX  
 CC The present sequence represents a Cytomegalovirus (CMV) minimal promoter  
 CC linked to 10 tet operator sequences. The specification describes a  
 CC method for regulating expression of a Tet (tetracycline) operator-linked  
 CC gene in a cell of a subject. The method comprises introducing into the  
 CC cell a nucleic acid encoding a fusion protein which inhibits  
 CC transcription in eukaryotic cells, the fusion protein comprising a  
 CC polypeptide which binds to a Tet operator sequence, operatively linked  
 CC to heterologous second polypeptide which inhibits transcription in  
 CC eukaryotic cells and modulating the concentration of a tetracycline  
 CC (analogue) in the subject. The method is used for the regulation of  
 CC gene expression system, using tetracycline (analogues). The system  
 CC enables a gene coupled to the system to be induced in the presence of  
 CC Tet and then stopped when Tet is removed.  
 ...  
 SQ Sequence 450 BP; 138 A; 101 C; 110 G; 101 T; 0 other;  
 Query Match 73.2%; Score 54.2; DB 19; Length 450;  
 Best Local Similarity 94.9%; Pred. No. 4.3e-12;  
 Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
 QY 2 GCGGTGTACGTTGGGAGGCCCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAACCGTC 60  
 DB 327 GCGGTGTACGTTGGGAGGCCCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAACCGTC 385  
 RESULT 11  
 AAV60081  
 ID AAV60081 standard; DNA; 450 BP.  
 AC AAV60081;  
 XX  
 XX 04-DEC-1998 (first entry)  
 DT  
 DE Cytomegalovirus minimal promoter linked to 10 tet operator sequences.  
 XX Tet repressor; tetracycline; regulation; expression;  
 KW Tet operator-linked gene; CMV minimal promoter; tet operator; ds.  
 XX Synthetic.  
 OS Human cytomegalovirus.  
 CC  
 US5814618-A.  
 XX 29-SEP-1998.  
 PD  
 XX 07-JUN-1995; 95US-0485978.  
 PF  
 XX 07-JUN-1995; 95US-0485978.  
 PR 14-JUN-1993; 93US-0076327.  
 PR 14-JUN-1993; 93US-0076327.  
 PR 14-JUN-1993; 93US-0076327.  
 PR 14-JUN-1993; 93US-0076327.  
 PR 01-JUL-1994; 94US-0270637.  
 PR 15-JUL-1994; 94US-0270637.  
 PR 06-FEB-1995; 95US-0383734.  
 XX (BADI ) BASF AG.  
 PA (KNOL ) KNOLL AG.  
 XX  
 PI Bujard H, Gossen M;  
 XX WPI; 1998-541795/46.  
 XX

PT Tetracycline based regulation of gene expression - uses a  
 PT tetracycline operator sequence joined to a gene of interest, the  
 PT gene of interest being induced in the presence, but not absence of  
 PT the antibiotic  
 XX  
 PS Disclosure; Columns 73-74; 63pp; English.  
 XX  
 CC The present sequence represents a Cytomegalovirus (CMV) minimal promoter  
 CC linked to 10 tet operator sequences. The specification describes a  
 CC method for regulating expression of a Tet (tetracycline) operator-linked  
 CC gene in a cell of a subject. The method comprises introducing into the  
 CC cell a nucleic acid encoding a fusion protein which inhibits  
 CC transcription in eukaryotic cells, the fusion protein comprising a  
 CC polypeptide which binds to a Tet operator sequence, operatively linked  
 CC to heterologous second polypeptide which inhibits transcription in  
 CC eukaryotic cells and modulating the concentration of a tetracycline  
 CC (analogue) in the subject. The method is used for the regulation of  
 CC gene expression system, using tetracycline (analogues). The system  
 CC enables a gene coupled to the system to be induced in the presence of  
 CC Tet and then stopped when Tet is removed.  
 XX  
 SQ Sequence 450 BP; 104 A; 109 C; 102 G; 135 T; 0 other;  
 Query Match 73.2%; Score 54.2; DB 19; Length 450;  
 Best Local Similarity 94.9%; Pred. No. 4.3e-12;  
 Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
 QY 2 GCGGTGTACGTTGGGAGGCCCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAACCGTC 60  
 DB 327 GCGGTGTACGTTGGGAGGCCCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAACCGTC 385  
 RESULT 12  
 AAX81721  
 ID AAX81721 standard; DNA; 450 BP.  
 AC AAX81721;  
 XX  
 XX 27-AUG-1999 (first entry)  
 DT  
 DE A human cytomegalovirus promoter sequence.  
 XX  
 XX Tetracycline-controllable transactivator; tTA; transgenic mouse;  
 KW transgene; gene function; gene expression; knockout animal;  
 KW Conditional inactivation; promoter; ds.  
 XX  
 OS Human cytomegalovirus.  
 XX  
 XX US5922927-A.  
 PN  
 XX 13-JUL-1999.  
 PD  
 XX 21-JUL-1997; 97US-0897719.  
 PF  
 XX 14-JUN-1994; 94US-0260452.  
 PR 14-JUN-1993; 93US-0076327.  
 PR 21-JUL-1997; 97US-0897719.  
 XX  
 XX (BADI ) BASF AG.  
 PA  
 XX Bujard H, Gossen M, Salfeld JG, Voss JW;  
 PI WPI; 1999-404496/34.  
 DR  
 XX Production of tetracycline-regulated transgenic mice, useful for  
 PT studying gene expression  
 PT  
 PS Example 2; Fig 6; 66pp; English.  
 XX  
 CC The present sequence represents a human cytomegalovirus promoter.  
 CC It is used to create the transgenic animals of the invention. The  
 CC specification describes a method for producing a transgenic mouse  
 CC having a transgene encoding a tetracycline-controllable transactivator

PR 01-JUL-1994; 94US-0270637.  
 PR 15-JUL-1994; 94US-0275876.  
 PR 03-FEB-1995; 95US-0383754.  
 XX  
 PA (BUJA/) BUJARD H.  
 PA (GOSS/) GOSSSEN M.  
 XX  
 PI Bujard H, Gossen M;  
 XX  
 DR WPI; 1996-087666/09.  
 XX  
 XX New tetracycline-regulated transcription modulators - comprising  
 PT fusion proteins which bind to tet operator sequences to activate or  
 PT inhibit transcription  
 XX  
 PS Disclosure; Page 73-74; 112pp; English.  
 XX  
 CC Fusion proteins comprising a first polypeptide which binds to a tet  
 CC operator sequence in the presence of tetracycline or a tetracycline  
 CC analogue, operatively linked to a second polypeptide which either  
 CC activates or inhibits transcription in eukaryotic cells may be used  
 CC to activate or inhibit transcription. Such proteins may be used to  
 CC regulate gene expression in cells and may be particularly useful for  
 CC gene therapy and for expression of gene products in transgenic  
 CC organisms. Induction of gene expression is rapid, efficient and  
 CC strong, typically 1000-2000 fold. The inducing agent does not cause  
 CC pleiotropic effects or cytotoxicity in eukaryotic cells.  
 CC Alternatively, a self regulating construct encoding a transactivator  
 CC fusion protein can be created. Here, the fusion protein is  
 CC operatively linked to a minimal promoter also comprising at least  
 CC one tet operator sequence. This sequence is a CMV minimal promoter  
 CC and ten tet operator sequences. Other promoters are described in  
 CC AAT11356 and AAT11357.  
 XX  
 SQ Sequence 450 BP; 138 A; 101 C; 110 G; 101 T; 0 other;  
 Query Match 73.2%; Score 54.2; DB 17; Length 450;  
 Best Local Similarity 94.9%; Pred. No. 4.3e-12;  
 Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
 OY 2 GCGGTGTACGTTGGAGCCCTATATAAGCAGAGCTCGTTTACTGAACCGTCAACCGTC 60  
 Db 327 GCGGTGTACGTTGGAGCCCTATATAAGCAGAGCTCGTTTACTGAACCGTCAACCGTC 385  
 RESULT 9  
 AAT11356  
 ID AAT11356 standard; DNA; 450 BP.  
 XX  
 AC AAT11356;  
 XX  
 DT 07-JUL-1996 (first entry)  
 XX  
 DE Minimal CMV promoter and ten tet operators.  
 XX  
 KW Tet repressor; Herpes simplex virus; HSV; virion protein 16; VP16;  
 KW fusion protein; gene expression; regulation; inhibition; activation;  
 KW transcription; ds.  
 XX  
 OS Transposon Tn-10/Cauliflower mosaic virus.  
 XX  
 PN WO9601313-A1.  
 XX  
 PD 18-JAN-1996.  
 XX  
 PF 29-JUN-1995; 95WO-US08179.  
 XX  
 PR 07-JUN-1995; 95US-0486814.  
 PR 01-JUL-1994; 94US-0270637.  
 PR 15-JUL-1994; 94US-0275876.  
 PR 03-FEB-1995; 95US-0383754.  
 XX  
 PA (BUJA/) BUJARD H.

PA (GOSS/) GOSSSEN M.  
 XX  
 PI Bujard H, Gossen M;  
 XX  
 DR WPI; 1996-087666/09.  
 XX  
 XX New tetracycline-regulated transcription modulators - comprising  
 PT fusion proteins which bind to tet operator sequences to activate or  
 PT inhibit transcription  
 XX  
 PS Disclosure; Page 74; 112pp; English.  
 XX  
 CC Fusion proteins comprising a first polypeptide which binds to a tet  
 CC operator sequence in the presence of tetracycline or a tetracycline  
 CC analogue, operatively linked to a second polypeptide which either  
 CC activates or inhibits transcription in eukaryotic cells may be used  
 CC to activate or inhibit transcription. Such proteins may be used to  
 CC regulate gene expression in cells and may be particularly useful for  
 CC gene therapy and for expression of gene products in transgenic  
 CC organisms. Induction of gene expression is rapid, efficient and  
 CC strong, typically 1000-2000 fold. The inducing agent does not cause  
 CC pleiotropic effects or cytotoxicity in eukaryotic cells.  
 CC Alternatively, a self regulating construct encoding a transactivator  
 CC fusion protein can be created. Here, the fusion protein is  
 CC operatively linked to a minimal promoter also comprising at least  
 CC one tet operator sequence. This sequence is a CMV minimal promoter  
 CC and ten tet operator sequences. Other promoters are described in  
 CC AAT11355 and AAT11357.  
 XX  
 SQ Sequence 450 BP; 104 A; 109 C; 102 G; 135 T; 0 other;  
 Query Match 73.2%; Score 54.2; DB 17; Length 450;  
 Best Local Similarity 94.9%; Pred. No. 4.3e-12;  
 Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
 OY 2 GCGGTGTACGTTGGAGCCCTATATAAGCAGAGCTCGTTTACTGAACCGTCAACCGTC 60  
 Db 327 GCGGTGTACGTTGGAGCCCTATATAAGCAGAGCTCGTTTACTGAACCGTCAACCGTC 385  
 RESULT 10  
 AAV60080  
 ID AAV60080 standard; DNA; 450 BP.  
 XX  
 AC AAV60080;  
 XX  
 DT 04-DEC-1998 (first entry)  
 XX  
 DE Cytomegalovirus minimal promoter linked to 10 tet operator sequences.  
 XX  
 KW Tet repressor; tetracycline; regulation; expression;  
 KW Tet operator-linked gene; CMV minimal promoter; tet operator; ds.  
 XX  
 OS Synthetic.  
 OS Human cytomegalovirus.  
 XX  
 PN US5814618-A.  
 XX  
 PD 29-SEP-1998.  
 XX  
 PF 07-JUN-1995; 95US-0485978.  
 XX  
 PR 07-JUN-1995; 95US-0485978.  
 PR 14-JUN-1993; 93US-0076327.  
 PR 14-JUN-1993; 93US-0076726.  
 PR 14-JUN-1994; 94US-0260452.  
 PR 01-JUL-1994; 94US-0270637.  
 PR 15-JUL-1994; 94US-0275876.  
 PR 06-FEB-1995; 95US-0383754.  
 XX  
 PA (BADI ) BASF AG.  
 PA (KNOL ) KNOLL AG.  
 XX

Qy	2	GGCGTGTACGGTGGGAGGCCTATATAAGCAGAGCTCGTTTACTGTAACCGTCAAACCGTC	60
Db	327	GGCGTGTACGGTGGGAGGCCTATATAAGCAGAGCTCGTTTACTGTAACCGTCAAACCGTC	385
RESULT 6			
AAQ76266			
ID	AAQ76266	standard; DNA; 450 BP.	
XX	AC	AC	
XX	AAQ76266;		
XX	XX		
DT	17-JUL-1995	(first entry)	
XX	XX		
DE	PhCMV*-1 promoter.		
XX	XX		
KW	tTA; transactivator; tetracycline-controllable transactivator;		
KW	conditional inactivation; homologous recombination; gene expression;		
KW	gene regulation; gene therapy; tetracycline-resistance; tetr;		
KW	transgenic animal; PhCMV*-1; promoter; tetr; CMV; ds.		
XX	XX		
OS	Human cytomegalovirus K12, Towne.		
XX	XX		
..			
Key	Location/Qualifiers		
mRNA	382..450		
	/*tag= a		
XX	XX		
PN	WO9429442-A.		
XX	XX		
PD	22-DEC-1994.		
XX	XX		
PF	14-JUN-1994; 94WO-US06734.		
XX	XX		
PR	14-JUN-1993; 93US-0076327.		
XX	XX		
PA	(BADI ) BASF AG.		
XX	XX		
PI	Bujard H, Gossen M, Salfeld JG, Voss JW;		
XX	XX		
DR	WPI; 1995-036472/05.		
XX	XX		
PT	Regulatory systems using tetracycline-controllable transactivator		
PT	(tTA) - useful for conditional inactivation or modulation of		
PT	gene expression in a host cell or animal		
XX	XX		
PS	Disclosure; Page 52-53; 103pp; English.		
CC	The minimal promoter PhCMV* spans the human CMV intermediate-		
CC	early promoter sequence from +75 to -53 (+1 being the first		
CC	nucleotide transcribed). TetO sequences were fused to		
CC	this core promoter to give the new promoters PhCMV*-1 (given		
CC	in AAQ76266) and PhCMV*-2 (AAQ76267). These promoters are used		
CC	to express tTA transactivator in host cells.		
XX	XX		
SQ	Sequence 450 BP; 138 A; 101 C; 110 G; 101 T; 0 other;		
Query Match	73.2%;	Score 54.2; DB 16; Length 450;	
Best Local Similarity	94.9%;	Pred. No. 4.3e-12;	
Matches	56; Conservative	0; Mismatches 3; Indels 0; Gaps	
Qy	2	GGCGTGTACGGTGGGAGGCCTATATAAGCAGAGCTCGTTTACTGTAACCGTCAAACCGTC	60
Db	327	GGCGTGTACGGTGGGAGGCCTATATAAGCAGAGCTCGTTTACTGTAACCGTCAAACCGTC	385
RESULT 7			
AAQ76267			
ID	AAQ76267	standard; DNA; 450 BP.	
XX	XX		
AC	AC		
XX	AAQ76267;		
XX	XX		
DT	17-JUL-1995	(first entry)	
XX	XX		
DE	PhCMV*-2 promoter.		

XX	TTA; transactivator; tetracycline-controllable transactivator;
KW	conditional inactivation; homologous recombination; gene expression;
KW	gene regulation; gene therapy; tetracycline-resistance; tetR;
KW	transgenic animal; PhCMV*-2; promoter; tetO; CMV; ds.
XX	
OS	Human cytomegalovirus Towne.
XX	
FH	Key
FT	Location/Qualifiers
FT	382..450
FT	/*tag= a
XX	
PN	WO9429442-A.
XX	
XX	22-DEC-1994.
XX	
PF	14-JUN-1994; 94WO-US06734.
XX	
PR	14-JUN-1993; 93US-0076327.
XX	
PA	(BADI ) BASF AG.
XX	
PI	Bujard H, Gossen M, Salfeld JG, Voss JW;
XX	
DR	WPI; 1995-036472/05.
XX	
PT	Regulatory systems using tetracycline-controllable transactivator
PT	(tTA) - useful for conditional inactivation or modulation of
PT	gene expression in a host cell or animal
XX	
PS	Disclosure; Page 53; 103pp; English.
XX	
CC	The minimal promoter PhCMV* spans the human CMV intermediate-
CC	early promoter sequence from +75 to -53 (+1 being the first
CC	nucleotide transcribed). TetO sequences were fused to
CC	this core promoter to give the new promoters PhCMV*-1 (given
CC	in AAQ76266) and PhCMV*-2 (AAQ76267). These promoters are used
CC	to express tTA transactivator in host cells.
XX	
SQ	Sequence 450 BP; 104 A; 109 C; 102 G; 135 T; 0 other;
	Query Match 73.2%; Score 54.2; DB 16; Length 450;
	Best Local Similarity 94.9%; Pred. No. 4.3e-12;
	Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps
Oy	2 GCGGTGTACGTGGGAGCCCTATATAAGCAGAGCTCGTTAGTGAACCGTCAACCGTC 60
Db	327 GCGGTGTACGTGGGAGCCCTATATAAGCAGAGCTCGTTAGTGAACCGTCAACCGTC 60
RESULT 8	
AA11355	
ID	AA11355 standard; DNA; 450 BP.
XX	
AC	AA11355;
XX	
DT	07-JUL-1996 (first entry)
XX	
DE	Minimal CMV promoter and ten tet operators.
XX	
KW	Tet repressor; Herpes simplex virus; HSV; virion protein 16; VP16;
KW	fusion protein; gene expression; regulation; inhibition; activation;
KW	transcription; ds.
XX	
OS	Transposon Tn-10/Cauliflower mosaic virus.
XX	
PN	WO9601313-A1.
XX	
PD	18-JAN-1996.
XX	
PF	29-JUN-1995; 95WO-US08179.
XX	
PR	07-JUN-1995; 95US-0486814.



XX Claim 12; Page 172; 241pp; English.

PS The invention relates to a construct which allows animals to be bred in

CC captivity but renders them infertile in the wild by allowing reversible

CC control over fertility and reproduction. The construct comprises a native

CC promoter, a blocking DNA sequence contoured for and designed to abrogate

CC a crucial gene's function or to cause its mis-expression, and a genetic

CC switch to regulate controlled expression/repression of the blocker/gene

CC knockout. The construct is useful for preventing embryogenesis or

CC gametogenesis in animals by stably transforming an animal cell with

CC the construct by microinjection, transfection or infection, where the

CC construct stably integrates into the genome by homologous recombination,

CC and implanting the cell into a host organism, where a whole animal

CC develops from the implanted cell. The present sequence is a repressible

CC promoter comprising tet responsive element (TRE) which is coupled to and

CC tightly regulates a minimal promoter region PminCMV from cytomegalovirus.

X

JQ Sequence 447 BP; 136 A; 104 C; 108 G; 99 T; 0 other;

Query Match 73.2%; Score 54.2; DB 22; Length 447;

Best Local Similarity 94.9%; Pred. No. 4.3e-12;

Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 GCGGTGTACGGTGGGAGCCCTATATAGCAGAGCTCGTTTACTGAACCGTCAACCGTC 60

|||||

Db 371 GCGGTGTACGGTGGGAGCCCTATATAGCAGAGCTCGTTTACTGAACCGTCAACCGTC 379

RESULT 4

AAT06869

ID AAT06869 standard; DNA; 450 BP.

XX AAT06869;

AC AAT06869;

XX 13-MAR-1996 (first entry)

XX PhCMV\*-1 tetO construct.

DE

DE PhCMV\*-1 tetO construct.

XX

XX Transactivator; tTA; tet operator; tetO; gene expression;

KW tetracycline-responsive promoter; PhCMV\*-1; human cytomegalovirus;

KW transcription activation; eukaryotic cell; ds; ss.

XX

OS Synthetic.

XX

XX Key Location/Qualifiers

FT TATA\_signal 347..353

FT /\*tag= a

XX

XX US5464758-A.

XX

XX 07-NOV-1995.

XX

XX 14-JUN-1993; 93US-0076726.

XX

XX 14-JUN-1993; 93US-0076726.

XX

XX (BUJA/) BUJARD H.

PA (GOSS/) GOSSSEN M.

XX

XX Bujard H, Gossen M;

PI

XX WPI; 1995-392612/50.

XX

XX Polynucleotide encoding trans:activator fusion protein contg. tet

PT repressor - used to control expression of gene regulated by minimal

PT promoter linked to tet operon, and vectors and cells where gene

PT expression is regulated by tetracycline

XX

PS Disclosure; Fig 6; 37pp; English.

XX

XX The PhCMV\*-1-tetO construct (AAT06869) is composed of tet operator

CC sequences separated by 95 bases from a human cytomegalovirus minimal

CC

CC IE promoter. A transactivator, tTA or tTAs (AAT06867-68), is used to

CC regulate expression of a heterologous gene operably linked to

CC PhCMV\*-1-tetO. On/off regulation of expression of the encoded

CC heterologous protein by host eukaryotic cells is provided by varying

CC the medium tetracycline conc.

XX

SQ Sequence 450 BP; 138 A; 101 C; 110 G; 101 T; 0 other;

Query Match 73.2%; Score 54.2; DB 16; Length 450;

Best Local Similarity 94.9%; Pred. No. 4.3e-12;

Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 GCGGTGTACGGTGGGAGCCCTATATAGCAGAGCTCGTTTACTGAACCGTCAACCGTC 60

|||||

Db 327 GCGGTGTACGGTGGGAGCCCTATATAGCAGAGCTCGTTTACTGAACCGTCAACCGTC 385

RESULT 5

AAT06870

ID AAT06870 standard; DNA; 450 BP.

XX AAT06870;

AC AAT06870;

XX 13-MAR-1996 (first entry)

XX

XX PhCMV\*-2 tetO construct.

DE

DE PhCMV\*-2 tetO construct.

XX

XX Transactivator; tTA; tet operator; tetO; gene expression;

KW tetracycline-responsive promoter; PhCMV\*-2; human cytomegalovirus;

KW transcription activation; eukaryotic cell; ds; ss.

XX

OS Synthetic.

XX

XX Key Location/Qualifiers

FT TATA\_signal 347..353

FT /\*tag= a

XX

XX US5464758-A.

XX

XX 07-NOV-1995.

XX

XX 14-JUN-1993; 93US-0076726.

XX

XX 14-JUN-1993; 93US-0076726.

XX

XX (BUJA/) BUJARD H.

PA (GOSS/) GOSSSEN M.

XX

XX Bujard H, Gossen M;

PI

XX WPI; 1995-392612/50.

XX

XX Polynucleotide encoding trans:activator fusion protein contg. tet

PT repressor - used to control expression of gene regulated by minimal

PT promoter linked to tet operon, and vectors and cells where gene

PT expression is regulated by tetracycline

XX

PS Disclosure; Fig 7; 37pp; English.

XX

XX The PhCMV\*-2-tetO construct (AAT06870) is composed of tet operator

CC sequences separated by 76 bases from a human cytomegalovirus minimal

CC IE promoter. A transactivator, tTA or tTAs (AAT06867-68), is used to

CC regulate expression of a heterologous gene operably linked to

CC PhCMV\*-2-tetO. On/off regulation of expression of the encoded

CC heterologous protein by host eukaryotic cells is provided by varying

CC the medium tetracycline conc.

XX

SQ Sequence 450 BP; 104 A; 109 C; 102 G; 135 T; 0 other;

Query Match 73.2%; Score 54.2; DB 16; Length 450;

Best Local Similarity 94.9%; Pred. No. 4.3e-12;

Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

PT mammalian gene expression, is based on the interferon regulating  
 PT factor-1 and its binding site  
 XX  
 PS Claim 1; Page 9; 19pp; English.  
 XX  
 CC The present invention describes a promoter-transactivator system for  
 CC inducible high-level expression of mammalian genes, and optionally  
 CC for control of cell growth. The promoter-transactivator system  
 CC comprises: (i) a promoter construct (IRFE promoter); and (ii)  
 CC transactivator construct encoding a fusion protein of IRF-1 (interferon  
 CC regulating factor-1) and the estrogen receptor (ER). The IRFE promoter  
 CC construct has the structure (MPSV-E)-(IRF-1 binding site)-(CMV)-DNA  
 CC where MPSV-E indicates the myeloproliferative sarcoma virus enhancer  
 CC repeat given in AAC64591; IRF-1 binding site given in AAC64592; and CMV  
 CC is the cytomegalovirus minimal promoter given in AAC64593 or their  
 CC functionally equivalent variants with one or more nucleotides  
 CC substituted, inserted or deleted. The promoter transactivator system is  
 CC a transcription regulator. Increased transcription results from binding  
 CC of the IRFE promoter to the transactivator, which is activated by  
 CC oestradiol or other ER ligands (these compounds displace the heat-shock  
 CC protein 90 which normally binds to the IRF-1/ER fusion, preventing its  
 CC activation). The system is used for increasing production of  
 CC therapeutically active proteins and where a IRF-1-green fluorescent  
 CC protein-human ER construct is used, for fluorescence-activated cell  
 CC sorting/analysis of transformed cells.  
 XX  
 SQ Sequence 74 BP; 19 A; 17 C; 23 G; 15 T; 0 other;

Query Match 100.0%; Score 74; DB 21; Length 74;  
 Best Local Similarity 100.0%; Pred. No. 2.5e-20;  
 Matches 74; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGGCGTGACGGTGGAGCGCTATATAGCAGAGCTCGTTAGTGAACCGTCAAAACCGTC 60  
 Db 1 TGGCGTGACGGTGGAGCGCTATATAGCAGAGCTCGTTAGTGAACCGTCAAAACCGTC 60

QY 61 AAACCGCGGAAGCT 74  
 Db 61 AAACCGCGGAAGCT 74

RESULT 2  
 AAT47525  
 ID AAT47525 standard; DNA; 432 BP.  
 XX  
 AC AAT47525;  
 XX  
 DT 08-AUG-1997 (first entry)  
 XX  
 DE tetO/ECMV chimeric target sequence encoding DNA.  
 YY Tetracycline; transactivator; anti-hepadnaviral agent; ss.

OS Chimeric - Human cytomegalovirus.  
 OS Chimeric - Escherichia coli.  
 XX  
 FH Key Location/Qualifiers  
 FT misc\_signal 1..312  
 FT /tag= a  
 FT /function= operator  
 FT /organism= Escherichia coli  
 FT promoter 313..432  
 FT /tag= b  
 FT /organism= Human cytomegalovirus

XX WO9639542-A1.  
 XX  
 XX 12-DEC-1996.  
 PD  
 XX 05-JUN-1996; 96WO-US08772.  
 PF  
 XX 05-JUN-1995; 95US-0462216.  
 PR  
 XX

PA (AVID-) AVID THERAPEUTICS.  
 XX  
 PI Barker CS; King RW, Seeger C;  
 XX  
 DR WPI; 1997-043163/04.  
 XX  
 PT Novel cell lines for inducing expression of hepatitis B virus genome  
 PT - used for cell-based assay to screen potential anti-hepadnaviral  
 PT agents  
 XX  
 PS Claim 8; Page 36; 49pp; English.  
 XX  
 CC A novel cell line capable of inducibly expressing a genome of  
 CC hepatitis B virus (HBV), is transformed with a HBV genome operably  
 CC linked to a target nucleotide sequence for activating expression of the  
 CC HBV genome, the activation of expression being caused by interaction of  
 CC the target sequence with a transactivator protein that specifically  
 CC interacts with the target sequence. The present sequence encodes a  
 CC target sequence which contains a prokaryotic tetracycline operator  
 CC (tetO) sequence fused to human cytomegalovirus promoter IE (hCMV).  
 CC The promoter is activated by the tetR/VP16 chimeric transactivator  
 CC protein (E.coli tet repressor fused to the activating domain of virion  
 CC protein 16 of herpes simplex virus). The method and cell lines can be  
 CC used for a cell-based assay to screen potential anti-hepadnaviral  
 CC agents.  
 XX  
 SQ Sequence 432 BP; 135 A; 94 C; 105 G; 98 T; 0 other;

Query Match 73.2%; Score 54.2; DB 18; Length 432;  
 Best Local Similarity 94.9%; Pred. No. 4.3e-12;  
 Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 GCGGTGACGGTGGAGCGCTATATAGCAGAGCTCGTTAGTGAACCGTCAAAACCGTC 60  
 Db 315 GCGGTGACGGTGGAGCGCTATATAGCAGAGCTCGTTAGTGAACCGTCAAGTCGCC 373

RESULT 3  
 AAD09962  
 ID AAD09962 standard; DNA; 447 BP.  
 XX  
 AC AAD09962;  
 XX  
 DT 12-SEP-2001 (first entry)  
 XX  
 DE Tet responsive element (TRE)-PminCMV repressible promoter.  
 XX  
 KW Tet responsive element; TRE; promoter; reproduction; embryogenesis; CMV;  
 KW cytomegalovirus; gametogenesis; microinjection; fertility; infection;  
 KW chimeric; ds.  
 XX  
 OS Chimeric - Cytomegalovirus.  
 OS Chimeric - Unidentified.  
 XX  
 PN WO200148224-A1.  
 XX  
 PD 05-JUL-2001.  
 XX  
 PF 22-DEC-2000; 2000WO-AU01596.  
 XX  
 PR 24-DEC-1999; 99AU-0004884.  
 XX  
 PA (CSIR ) COMMONWEALTH SCI & IND RES ORG.  
 XX  
 PI Thresher R, Hinds L, Hardy C, Whyard S, Vignarajan S, Grewe PM;  
 PI Patil J;  
 XX  
 DR WPI; 2001-425672/45.  
 XX  
 PT Novel construct for preventing embryogenesis in animals comprises  
 PT native promoter, blocking DNA which abrogates function of crucial gene  
 PT and genetic switch to regulate expression/repression of blocker/gene  
 PT knockout -

GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: March 5, 2003, 22:29:12 ; Search time 72.2126 Seconds

(without alignments)

2307.741 Million cell updates/sec

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Gapop 10.0 , Gapext 1.0

searched: 2185239 seqs, 1125999159 residues

Total number of hits satisfying chosen parameters: 4370478

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Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	74	100.0	74	AAC64593	Cytomegalovirus mi
2	54.2	73.2	432	AAAT47525	tetO/cwM chimeric
3	54.2	73.2	447	AAAD09962	tet responsive ele
4	54.2	73.2	450	AAAT06869	PhCMV*-1 tetO cons
5	54.2	73.2	450	AAAT06870	PhCMV*-2 tetO cons
6	54.2	73.2	450	AAQ76266	PhCMV*-1 promoter.
7	54.2	73.2	450	AAQ76267	PhCMV*-2 promoter.
8	54.2	73.2	450	AAAT1355	Minimal CMV promot
9	54.2	73.2	450	AAAT1356	Minimal CMV promot

10	54.2	73.2	450	19	AAV60080	Cytomegalovirus mi
11	54.2	73.2	450	19	AAV60081	Cytomegalovirus mi
12	54.2	73.2	450	20	AAAX81721	A human cytomegalo
13	54.2	73.2	450	20	AAAX81722	A human cytomegalo
14	54.2	73.2	450	20	AAAX60046	Minimal Human cyto
15	54.2	73.2	450	20	AAAX60047	Minimal Human cyto
16	54.2	73.2	450	20	AAAX27901	PhCMV*-1 promoter
17	54.2	73.2	450	20	AAAX27903	PhCMV*-2 promoter
18	54.2	73.2	450	20	AAAX01365	PhCMV*-1 sequence.
19	54.2	73.2	450	20	AAAX01367	PhCMV*-2 sequence.
20	54.2	73.2	450	21	AAZ56127	Tetracycline respo
21	54.2	73.2	450	21	AAZ56128	Tetracycline respo
22	54.2	73.2	450	22	AAAD09834	Human cytomegalovi
23	54.2	73.2	450	22	AAAD09835	Human cytomegalovi
24	54.2	73.2	450	22	AAH25573	Nucleotide sequenc
25	54.2	73.2	450	22	AAH25574	Nucleotide sequenc
26	54.2	73.2	450	22	AAH47632	Tetracycline-respo
27	54.2	73.2	450	22	AAH47633	Tetracycline-respo
28	54.2	73.2	470	20	AAAX90500	Nucleotide sequenc
29	54.2	73.2	481	19	AAV17755	Tet promoter clone
30	54.2	73.2	481	19	AAV18691	Inducible promoter
31	54.2	73.2	500	24	ABL41162	Cytomegalovirus im
32	54.2	73.2	520	17	AAAT11359	teto/minimal CMV t
33	54.2	73.2	520	18	AAAT45722	Bidirectional prom
34	54.2	73.2	520	19	AAV60079	Bi-directional tet
35	54.2	73.2	520	20	AAAX60045	Bidirectional prom
36	54.2	73.2	520	21	AAZ56126	Bidirectional prom
37	54.2	73.2	520	22	AAH25572	Nucleotide sequenc
38	54.2	73.2	520	22	AAH47631	Nucleotide sequenc
39	54.2	73.2	569	17	AAAT11358	Bidirectional prom
40	54.2	73.2	569	18	AAAT45721	Bi-directional tet
41	54.2	73.2	569	19	AAV60078	Bidirectional prom
42	54.2	73.2	569	20	AAAX60044	Bidirectional prom
43	54.2	73.2	569	21	AAZ56125	Bidirectional prom
44	54.2	73.2	569	22	AAH25571	Nucleotide sequenc
45	54.2	73.2	569	22	AAH47630	Nucleotide sequenc

#### ALIGNMENTS

RESULT 1  
AAC64593  
ID AAC64593 standard; DNA; 74 BP.  
XX  
AC AAC64593;  
XX  
DT 15-FEB-2001 (first entry)  
XX  
DE Cytomegalovirus minimal promoter sequence SEQ ID NO:1.  
XX  
KW Myeloproliferative sarcoma virus; MPSV; MPSV-E; enhancer; CMV;  
KW Cytomegalovirus; firefly; IRF-1 binding site; minimal promoter;  
KW interferon regulatory factor 1 binding site; transcription regulator;  
KW promoter transactivator system; ds.  
XX  
OS Cytomegalovirus.  
XX  
PN EP1046710-AL.  
XX  
PD 25-OCT-2000.  
XX  
PF 23-APR-1999; 99EP-0108068.  
XX  
PR 23-APR-1999; 99EP-0108068.  
XX  
PA (GBFB ) GES BIOTECHNOLOGISCHE FORSCHUNG MBH.  
XX  
PI Mueller PP, Geserick C, Schroeder K, Hauser H;  
XX WPI; 2000-648930/63.  
XX  
PT Promoter-transactivator system, useful for inducing high level

VERSION AR072128.1 GI:7223016  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 450)  
AUTHORS Bujard,H. and Gossen,M.  
TITLE Mice transgenic for a tetracycline-inducible transcriptional activator  
JOURNAL Patent: US 5912411-A 8 15-JUN-1999;  
FEATURES Location/Qualifiers  
source 1..450  
BASE COUNT 138 a 101 c 110 g 101 t  
ORIGIN

Query Match 73.2%; Score 54.2; DB 6; Length 450;  
Best Local Similarity 94.9%; Pred. No. 8.5e-11;  
Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 2 GCGGTGACGGTGGAGCGCTATATAAGCAGAGCTCGTTAGTGAACCGTCAAAACCGTC 60  
|||||  
327 GCGGTGACGGTGGAGCGCTATATAAGCAGAGCTCGTTAGTGAACCGTCAATCGCC 385  
|||||

RESULT 12  
AR072129  
LOCUS AR072129 450 bp DNA linear PAT 18-FEB-2000  
DEFINITION Sequence 9 from patent US 5912411.  
ACCESSION AR072129  
VERSION AR072129.1 GI:7223017  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 450)  
AUTHORS Bujard,H. and Gossen,M.  
TITLE Mice transgenic for a tetracycline-inducible transcriptional activator  
JOURNAL Patent: US 5912411-A 9 15-JUN-1999;  
FEATURES Location/Qualifiers  
source 1..450  
BASE COUNT 104 a 109 c 102 g 135 t  
ORIGIN

Query Match 73.2%; Score 54.2; DB 6; Length 450;  
Best Local Similarity 94.9%; Pred. No. 8.5e-11;  
Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 2 GCGGTGACGGTGGAGCGCTATATAAGCAGAGCTCGTTAGTGAACCGTCAAAACCGTC 60  
|||||  
327 GCGGTGACGGTGGAGCGCTATATAAGCAGAGCTCGTTAGTGAACCGTCAATCGCC 385  
|||||

RESULT 13  
AR095982  
LOCUS AR095982 450 bp DNA linear PAT 08-SEP-2000  
DEFINITION Sequence 8 from patent US 6004941.  
ACCESSION AR095982  
VERSION AR095982.1 GI:10024370  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 450)  
AUTHORS Bujard,H. and Gossen,M.  
TITLE Methods for regulating gene expression  
JOURNAL Patent: US 6004941-A 8 21-DEC-1999;  
FEATURES Location/Qualifiers  
source 1..450  
BASE COUNT 138 a 101 c 110 g 101 t  
ORIGIN

Query Match 73.2%; Score 54.2; DB 6; Length 450;  
Best Local Similarity 94.9%; Pred. No. 8.5e-11;  
Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 2 GCGGTGACGGTGGAGCGCTATATAAGCAGAGCTCGTTAGTGAACCGTCAAAACCGTC 60  
|||||  
327 GCGGTGACGGTGGAGCGCTATATAAGCAGAGCTCGTTAGTGAACCGTCAATCGCC 385  
|||||

RESULT 14  
AR095983  
LOCUS AR095983 450 bp DNA linear PAT 08-SEP-2000  
DEFINITION Sequence 9 from patent US 6004941.  
ACCESSION AR095983  
VERSION AR095983.1 GI:10024372  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 450)  
AUTHORS Bujard,H. and Gossen,M.  
TITLE Methods for regulating gene expression  
JOURNAL Patent: US 6004941-A 9 21-DEC-1999;  
FEATURES Location/Qualifiers  
source 1..450  
BASE COUNT 104 a 109 c 102 g 135 t  
ORIGIN

Query Match 73.2%; Score 54.2; DB 6; Length 450;  
Best Local Similarity 94.9%; Pred. No. 8.5e-11;  
Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 2 GCGGTGACGGTGGAGCGCTATATAAGCAGAGCTCGTTAGTGAACCGTCAAAACCGTC 60  
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327 GCGGTGACGGTGGAGCGCTATATAAGCAGAGCTCGTTAGTGAACCGTCAATCGCC 385  
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RESULT 15  
AR136642  
LOCUS AR136642 450 bp DNA linear PAT 16-JUN-2001  
DEFINITION Sequence 8 from patent US 6136954.  
ACCESSION AR136642  
VERSION AR136642.1 GI:14477314  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 450)  
AUTHORS Bujard,H. and Gossen,M.  
TITLE Tetracycline-inducible transcriptional activator fusion proteins  
JOURNAL Patent: US 6136954-A 8 24-OCT-2000;  
FEATURES Location/Qualifiers  
source 1..450  
BASE COUNT 138 a 101 c 110 g 101 t  
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Query Match 73.2%; Score 54.2; DB 6; Length 450;  
Best Local Similarity 94.9%; Pred. No. 8.5e-11;  
Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 2 GCGGTGACGGTGGAGCGCTATATAAGCAGAGCTCGTTAGTGAACCGTCAAAACCGTC 60  
|||||  
327 GCGGTGACGGTGGAGCGCTATATAAGCAGAGCTCGTTAGTGAACCGTCAATCGCC 385  
|||||

Search completed: March 5, 2003, 23:34:39  
Job time : 408.715 secs

SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 450)  
AUTHORS Bujard,H. and Gossen,M., Salfield,J.G. and Voss,J.W.  
TITLE Mice transgenic for a tetracycline-controlled transcriptional activator  
JOURNAL Patent: US 5859310-A 6 12-JAN-1999;  
FEATURES Location/Qualifiers  
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          /organism="unknown"  
BASE COUNT 104 a 109 c 102 g 135 t  
ORIGIN

Query Match 73.2%; Score 54.2; DB 6; Length 450;  
Best Local Similarity 94.9%; Pred. No. 8.5e-11;  
Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Y 2 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTACTGAACCGTCAAAACCGTC 60  
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Db 327 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTACTGAACCGTCAAGTCGCC 385

RESULT 7  
AR032151  
LOCUS AR032151 450 bp DNA linear PAT 29-SEP-1999  
DEFINITION Sequence 8 from patent US 5866755.  
ACCESSION AR032151  
VERSION AR032151.1 GI:5946440  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 450)  
AUTHORS Bujard,H. and Gossen,M.  
TITLE Animals transgenic for a tetracycline-regulated transcriptional inhibitor  
JOURNAL Patent: US 5866755-A 8 02-FEB-1999;  
FEATURES Location/Qualifiers  
          1..450  
          /organism="unknown"  
BASE COUNT 138 a 101 c 110 g 101 t  
ORIGIN

Query Match 73.2%; Score 54.2; DB 6; Length 450;  
Best Local Similarity 94.9%; Pred. No. 8.5e-11;  
Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Y 2 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTACTGAACCGTCAAAACCGTC 60  
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Db 327 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTACTGAACCGTCAAGTCGCC 385

RESULT 8  
AR032152  
LOCUS AR032152 450 bp DNA linear PAT 29-SEP-1999  
DEFINITION Sequence 9 from patent US 5866755.  
ACCESSION AR032152  
VERSION AR032152.1 GI:5946441  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 450)  
AUTHORS Bujard,H. and Gossen,M.  
TITLE Animals transgenic for a tetracycline-regulated transcriptional inhibitor  
JOURNAL Patent: US 5866755-A 9 02-FEB-1999;  
FEATURES Location/Qualifiers  
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          /organism="unknown"  
BASE COUNT 104 a 109 c 102 g 135 t  
ORIGIN

Query Match 73.2%; Score 54.2; DB 6; Length 450;  
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Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Y 2 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTACTGAACCGTCAAAACCGTC 60  
|||||  
Db 327 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTACTGAACCGTCAAGTCGCC 385

RESULT 9  
AR043816  
LOCUS AR043816 450 bp DNA linear PAT 29-SEP-1999  
DEFINITION Sequence 8 from patent US 5814618.  
ACCESSION AR043816  
VERSION AR043816.1 GI:5964824  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 450)  
AUTHORS Bujard,H. and Gossen,M.  
TITLE Methods for regulating gene expression  
JOURNAL Patent: US 5814618-A 8 29-SEP-1998;  
FEATURES Location/Qualifiers  
          1..450  
          /organism="unknown"  
BASE COUNT 138 a 101 c 110 g 101 t  
ORIGIN

Query Match 73.2%; Score 54.2; DB 6; Length 450;  
Best Local Similarity 94.9%; Pred. No. 8.5e-11;  
Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Y 2 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTACTGAACCGTCAAAACCGTC 60  
|||||  
Db 327 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTACTGAACCGTCAAGTCGCC 385

RESULT 10  
AR043817  
LOCUS AR043817 450 bp DNA linear PAT 29-SEP-1999  
DEFINITION Sequence 9 from patent US 5814618.  
ACCESSION AR043817  
VERSION AR043817.1 GI:5964825  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 450)  
AUTHORS Bujard,H. and Gossen,M.  
TITLE Methods for regulating gene expression  
JOURNAL Patent: US 5814618-A 9 29-SEP-1998;  
FEATURES Location/Qualifiers  
          1..450  
          /organism="unknown"  
BASE COUNT 104 a 109 c 102 g 135 t  
ORIGIN

Query Match 73.2%; Score 54.2; DB 6; Length 450;  
Best Local Similarity 94.9%; Pred. No. 8.5e-11;  
Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Y 2 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTACTGAACCGTCAAAACCGTC 60  
|||||  
Db 327 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTACTGAACCGTCAAGTCGCC 385

RESULT 11  
AR072128  
LOCUS AR072128 450 bp DNA linear PAT 18-FEB-2000  
DEFINITION Sequence 8 from patent US 5912411.  
ACCESSION AR072128

FEATURES Gesellschaft fjr Biotechnologische Forschung mbH (GBF) (: DB)

source 1..74  
Location/Qualifiers  
/organism="unidentified"  
/db\_xref="taxon:32644"  
BASE COUNT 19 a 17 c 23 g 15 t  
ORIGIN

Query Match 100.0%; Score 74; DB 6; Length 74;  
Best Local Similarity 100.0%; Pred. No. 7.5e-19;  
Matches 74; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGGCGGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAAAACCGTC 60  
|||||  
Db 1 TGGCGGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAAAACCGTC 60

Qy 61 AAACCGGGAAGCT 74  
|||||

Db 61 AAACCGGGAAGCT 74

RESULT 2  
US 18986  
LOCUS AR020199 432 bp DNA linear PAT 05-DEC-1998  
DEFINITION Sequence 1 from patent US 5789156.  
ACCESSION AR020199  
VERSION AR020199.1 GI:3974814  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 432)  
AUTHORS King,R.W., Barker,C.S. and Seeger,C.  
TITLE Cultured cell line that inducibly expresses the hepatitis B virus genome, and uses thereof for screening antiviral substances  
JOURNAL Patent: US 5723319-A 1 03-MAR-1998;  
FEATURES Location/Qualifiers  
source 1..432  
/organism="unknown"

BASE COUNT 135 a 94 c 105 g 98 t  
ORIGIN

Query Match 73.2%; Score 54.2; DB 6; Length 432;  
Best Local Similarity 94.9%; Pred. No. 8.5e-11;  
Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 2 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAAAACCGTC 60  
|||||  
Db 315 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAAGTCGCC 373

ULT 3  
J20199  
LOCUS AR020199 450 bp DNA linear PAT 05-DEC-1998  
DEFINITION Sequence 8 from patent US 5789156.  
ACCESSION AR020199  
VERSION AR020199.1 GI:3974814  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 450)  
AUTHORS Bujard,H. and Gossen,M.  
TITLE Tetracycline-regulated transcriptional inhibitors  
JOURNAL Patent: US 5789156-A 8 04-AUG-1998;  
FEATURES Location/Qualifiers  
source 1..450  
/organism="unknown"

BASE COUNT 138 a 101 c 110 g 101 t  
ORIGIN

Query Match 73.2%; Score 54.2; DB 6; Length 450;  
Best Local Similarity 94.9%; Pred. No. 8.5e-11;

Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
Qy 2 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAAAACCGTC 60  
|||||  
Db 327 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAAGTCGCC 385

RESULT 4  
LOCUS AR020200 450 bp DNA linear PAT 05-DEC-1998  
DEFINITION Sequence 9 from patent US 5789156.  
ACCESSION AR020200  
VERSION AR020200.1 GI:3974815  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 450)  
AUTHORS Bujard,H. and Gossen,M.  
TITLE Tetracycline-regulated transcriptional inhibitors  
JOURNAL Patent: US 5789156-A 9 04-AUG-1998;  
FEATURES Location/Qualifiers  
source 1..450  
/organism="unknown"

BASE COUNT 104 a 109 c 102 g 135 t  
ORIGIN

Query Match 73.2%; Score 54.2; DB 6; Length 450;  
Best Local Similarity 94.9%; Pred. No. 8.5e-11;  
Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 2 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAAAACCGTC 60  
|||||  
Db 327 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAAGTCGCC 385

RESULT 5  
LOCUS AR029414 450 bp DNA linear PAT 29-SEP-1999  
DEFINITION Sequence 5 from patent US 5859310.  
ACCESSION AR029414  
VERSION AR029414.1 GI:5941387  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 450)  
AUTHORS Bujard,H., Gossen,M., Salfeld,J.G. and Voss,J.W.  
TITLE Mice transgenic for a tetracycline-controlled transcriptional activator  
JOURNAL Patent: US 5859310-A 5 12-JAN-1999;  
FEATURES Location/Qualifiers  
source 1..450  
/organism="unknown"

BASE COUNT 138 a 101 c 110 g 101 t  
ORIGIN

Query Match 73.2%; Score 54.2; DB 6; Length 450;  
Best Local Similarity 94.9%; Pred. No. 8.5e-11;  
Matches 56; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 2 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAAAACCGTC 60  
|||||  
Db 327 GCGGTGTACGGTGGGAGCCCTATATAAGCAGAGCTCGTTTGTAGTGAACCGTCAAGTCGCC 385

RESULT 6  
LOCUS AR029415 450 bp DNA linear PAT 29-SEP-1999  
DEFINITION Sequence 6 from patent US 5859310.  
ACCESSION AR029415  
VERSION AR029415.1 GI:5941388  
KEYWORDS

GenCore version 5.1.3  
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model  
Run on: March 5, 2003, 22:29:16 ; Search time 407.715 Seconds  
(without alignments)  
5282.140 Million cell updates/sec  
Title: US-09-980-548-3  
Perfect score: 74  
Sequence: 1 tggcgtgtacgtgtggagc.....accgtcaaacgcggaagct 74

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0  
searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0  
Maximum DB seq length: 20000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

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2:	gb_htg:	*	2	54.2	73.2	432	6	I89986	Sequence 1
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4:	gb_om:	*	4	54.2	73.2	450	6	AR020200	Sequence 1
5:	gb_ov:	*	5	54.2	73.2	450	6	AR029414	Sequence 1
6:	gb_pat:	*	6	54.2	73.2	450	6	AR029415	Sequence 1
7:	gb_ph:	*	7	54.2	73.2	450	6	AR032151	Sequence 1
8:	gb_pl:	*	8	54.2	73.2	450	6	AR032152	Sequence 1
9:	gb_pr:	*	9	54.2	73.2	450	6	AR043816	Sequence 1
10:	gb_ro:	*	10	54.2	73.2	450	6	AR043817	Sequence 1
11:	gb_sts:	*	11	54.2	73.2	450	6	AR072128	Sequence 1
12:	gb_sy:	*	12	54.2	73.2	450	6	AR072129	Sequence 1
13:	gb_un:	*	13	54.2	73.2	450	6	AR095982	Sequence 1
14:	gb_vl:	*	14	54.2	73.2	450	6	AR095983	Sequence 1
15:	em_ba:	*	15	54.2	73.2	450	6	AR136642	Sequence 1
16:	em_fun:	*	16	54.2	73.2	450	6	AR136643	Sequence 1
17:	em_in:	*	17	54.2	73.2	450	6	AR157177	Sequence 1
18:	em_mu:	*	18	54.2	73.2	450	6	AR157178	Sequence 1
19:	em_mu:	*	19	54.2	73.2	450	6	AR164126	Sequence 1
20:	em_mu:	*	20	54.2	73.2	450	6	AR164127	Sequence 1
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22:	em_mu:	*	22	54.2	73.2	450	6	AX127250	Sequence 1
23:	em_mu:	*	23	54.2	73.2	450	6	I15364	Sequence 12
24:	em_mu:	*	24	54.2	73.2	450	6	I15365	Sequence 13
25:	em_mu:	*	25	54.2	73.2	450	6	I32812	Sequence 8
26:	em_mu:	*	26	54.2	73.2	450	6	I32813	Sequence 9
27:	em_mu:	*	27	54.2	73.2	450	6	I56756	Sequence 5
28:	em_mu:	*	28	54.2	73.2	450	6	I56757	Sequence 6
29:	em_mu:	*	29	54.2	73.2	450	6	I59626	Sequence 8
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34:	em_mu:	*	34	54.2	73.2	520	6	AR032150	Sequence 1
35:	em_mu:	*	35	54.2	73.2	520	6	AR043815	Sequence 1
36:	em_mu:	*	36	54.2	73.2	520	6	AR072127	Sequence 1
37:	em_mu:	*	37	54.2	73.2	520	6	AR095981	Sequence 1
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39:	em_mu:	*	39	54.2	73.2	520	6	AR157176	Sequence 1
40:	em_mu:	*	40	54.2	73.2	520	6	AR164125	Sequence 1
41:	em_mu:	*	41	54.2	73.2	520	6	AX127248	Sequence 1
42:	em_mu:	*	42	54.2	73.2	520	6	I32811	Sequence 7
43:	em_mu:	*	43	54.2	73.2	520	6	I59625	Sequence 7
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45:	em_mu:	*	45	54.2	73.2	569	6	AR032149	Sequence 1

ALIGNMENTS

RESULT 1  
AX040911  
LOCUS  
DEFINITION  
Accession  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL

AX040911  
Sequence 3 from Patent WO0065074.  
AX040911  
AX040911.1 GI:11340533  
unidentified.  
unidentified.  
unclassified.  
1 (bases 1 to 74)  
Mueller.P., Geserick.C., Schroeder.K. and Hauser.H.  
Promoter-transactivator system for inducible high-level mammalian  
gene expression with the option of cell growth control  
Patent: WO 0065074-A 3 02-NOV-2000;

74 bp  
DNA  
linear  
PAT 23-NOV-2000

Pred. No. is the number of results predicted by chance to have a





VERSION BI938198.1 GI:16252670



[illegible]



adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (gII4732114[gblAF129072.1], a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent *E. coli* XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT	127 a	130 c	133 g	127 t
ORIGIN				
Query Match		42.3%	Score 29.6;	DB 17; Length 517;
Best Local Similarity		88.9%;	Pred. No. 30;	
Matches 32;	Conservative	0;	Mismatches	4; Indels 0; Gaps 0;

QY 4 CCCTTCTCGGAAATGGAAACTGAAAAATCAGATCCC 39  
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 Db 309 CCCTTCTCGGAAATGGAAACTGAAAAATCTACTTCC 274

RESULT 5	BI936879/c	LOCUS	BI936879	391 bp	mRNA	linear	EST 18-OCT-2001
DEFINITION			dd16b07.y1 Wellcome CRC pSK St.10.5 <i>Xenopus laevis</i> cdNA clone				
			IMAGE:3420612 5' similar to TR:014597 014597 NON-FUNCTIONAL FOLATE BINDING PROTEIN. ; mRNA sequence.				

ACCESSION BI936879  
VERSION BI936879.1 GI:16251351

KEYWORDS EST.  
SOURCE African clawed frog.  
ORGANISM *Xenopus laevis*  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Amphibia; Batrachia; Anura; Mesobatrachia; Pipidoidea; Pipidae;  
Xenopodinae; *Xenopus*.

REFERENCE  
1 (bases 1 to 391)

AUTHORS  
Clifton, S., Johnson, S.L., Blumberg, B., Song, J., Hillier, L., Pape, D., Martin, J., Wylie, T., Underwood, K., Theising, B., Bowers, Y., Person, B., Gibbons, M., Harvey, N., Ritter, E., Jackson, Y., McCann, R., Waterston, R. and Wilson, R.

**TITLE**  
**Washu Xenopus EST project, 1999**

TITLE	WasNu Xenopus EST project, 1999
JOURNAL	Unpublished (1999)
COMMENT	Contact: Sandy Clifton, ph.d. WasNu Xenopus EST project, 1999 Washington University School of Medicine 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA Tel: 314 286 1800 Fax: 314 286 1810

Tel: 314 286 1800  
Fax: 314 286 1810  
Email: estewatson.wustl.edu

Library constructed by N. Garrett, P. Lemaire, A. M. Zorn, and J.B. Gordon (Wellcome/CRC Institute). DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: Xenopus clones from this library are available through the I.M.A.G.E. Consortium/LLNL at: [infoimage.llnl.gov](mailto:infoimage.llnl.gov)

Possible reversed clone: similarity on wrong strand

Seq primer: -40RP from Gibco

High quality sequence stop: 165.

```

Possible reversed clone: similarity on wrong strand
Seq primer: -40RP from Gibco
High quality sequence stop: 165.
Location/Qualifiers
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        /db_xref="taxon:8355"
        /clone="IMAGE:3420612"
        /clone_lib="Wellcome CRC psk st 10 5"
        /tissue_type="embryo, stage 10.5"
        /lab_host="DH10B (phage-resistant)"
        /note="Vector: pBluescript SK; Site_1: NotI; Site_2:
        EcoRI; cDNAs were oligo-dT primed and directionally
        cloned. Staging according to Nieuwkoop and Faber. Library
        was constructed by N. Garrett, P. Lemaire, A.M. Zorn, and
        J.B. Gurdon (Wellcome/CRC Institute). Note: This is a
        xenopus Gene Collection (XGC) library."
88 a 106 c 90 g 107 t
BASE COUNT

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GenCore version 5.1.3  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: March 5, 2003, 22:43:11 ; Search time 527.874 Seconds  
(without alignments)  
2147.640 Million cell updates/sec

Title: US-09-980-548-2  
Perfect score: 70  
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Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Jearched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000  
Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : EST.\*

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5: em\_estov:\*  
6: em\_estpl:\*  
7: em\_estro:\*  
8: em\_htc:\*  
9: gb\_est1:\*  
10: gb\_est2:\*  
11: gb\_htc:\*  
12: gb\_est3:\*  
13: gb\_est4:\*  
14: gb\_est5:\*  
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

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5	28.2	40.3	391	13	BI936879
6	28.2	40.3	540	17	AZ913831

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	20	27	38.6	695	17	AZ812128
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	22	26.8	38.3	524	13	BU032573
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c	43	26.2	37.4	603	13	BI503827
c	44	26.2	37.4	668	13	BJ071703
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## ALIGNMENTS

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LOCUS	RC1-CT0249-130900-212-b06	CT0249	Homo sapiens	CDNA	mRNA sequence.
DEFINITION	BQ321577				
ACCESSION	BQ321577.1	GI:20929928			
VERSION	EST.				
KEYWORDS	human.				
SOURCE	Homo sapiens				
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.				
REFERENCE	1 (bases 1 to 506)				
AUTHORS	Dias Neto,B., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R., Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Cosca,F.F., Goldman,G.H., Carvalho,A.F., Matsumura,A., Baia,G.S., Simpson,D.H., Brunstein,A., deOliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and Simpson,A.J.				
TITLE	Shotgun sequencing of the human transcriptome with ORF expressed sequence tags				
JOURNAL	Proc. Natl. Acad. Sci. U.S.A.	97 (7),	3491-3496	(2000)	
MEDLINE	20202663				
COMMENT	Contact: Simpson A.J.G. Laboratory of Cancer Genetics Ludwig Institute for Cancer Research Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil Tel: +55-11-2704922 Fax: +55-11-2707001				

Thu Mar 6 09:56:00 2003

us-09-980-548-2.rng

Page 12

Search completed: March 5, 2003, 22:49:56  
Job time : 73.3092 secs



PS Claim 25; SEQ ID No 4962; 487pp; English.  
XX  
CC The present invention relates to human single exon nucleic acid probes  
CC (SENPs). The present sequence is one such probe. The SENPs are derived  
CC from human Hela cells. The SENPs can be used to produce a single exon  
CC microarray, which can be used for measuring human gene expression in a  
CC sample derived from human cervical epithelial cells. By measuring gene  
CC expression, the probes are therefore useful in grading and/or staging  
CC of diseases of the cervix, notably cervical cancer.  
CC Note: The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences.  
XX  
SQ Sequence 415 BP; 136 A; 91 C; 80 G; 108 T; 0 other;  
  
Query Match 36.6%; Score 25.6; DB 22; Length 415;  
Best Local Similarity 66.1%; Pred. No. 23;  
Matches 37; Conservative 0; Mismatches 19; Indels 0; Gaps 0;  
  
QY 15 AAATGGAACCTGAAATCAGATCCCTTCGCGGAATGGAACCTGAAATCAGATC 70  
IIII II III III III III I II III III III III II  
Db 201 AAATGACATAAAATTCACATCCATGACCTAAATGGAATGGAATCAGATC 256  
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ID AAI36372 standard; DNA; 415 BP.  
XX  
AC AAI36372;  
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XX  
DT 17-OCT-2001 (first entry)  
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DE Probe #5058 used to measure gene expression in human placenta sample.  
XX  
KW Probe; microarray; human; placenta; antenatal diagnosis;  
KW genetic disorder; ss.  
XX  
OS Homo sapiens.  
XX  
XX  
PN WO200157272-A2.  
XX  
PD 09-AUG-2001.  
XX  
PF 30-JAN-2001; 2001WO-US00663.  
XX  
PS 04-FEB-2000; 2000US-0180312.  
XX 26-MAY-2000; 2000US-0207456.  
XX 30-JUN-2000; 2000US-0608408.  
XX 03-AUG-2000; 2000US-0632366.  
XX 21-SEP-2000; 2000US-0234687.  
XX 27-SEP-2000; 2000US-0236359.  
XX 04-OCT-2000; 2000GB-0024263.  
XX  
PA (MOLE-) MOLECULAR DYNAMICS INC.  
XX  
XX  
PI Penn SG, Hanzel DK, Chen W, Rank DR;  
XX WO200157272-A2.  
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PD 09-AUG-2001.  
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PS 04-FEB-2000; 2000US-0180312.  
XX 26-MAY-2000; 2000US-0207456.  
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XX 21-SEP-2000; 2000US-0234687.  
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XX 04-OCT-2000; 2000GB-0024263.  
XX  
PA (MOLE-) MOLECULAR DYNAMICS INC.  
XX  
XX  
PI Penn SG, Hanzel DK, Chen W, Rank DR;  
XX WPI; 2001-488897/53.  
XX  
DR Human genome-derived single exon nucleic acid probes useful for  
PT analyzing gene expression in human placenta -  
XX  
XX  
PS Claim 25; SEQ ID No 5058; 654pp; English.  
XX  
XX The present invention relates to single exon nucleic acid probes (SENPs).  
CC The present sequence is one such probe. The probes are useful for  
CC producing a microarray for predicting, measuring and displaying gene  
CC expression in samples derived from human placenta. The probes are useful  
CC for antenatal diagnosis of human genetic disorders.  
XX  
XX  
SQ Sequence 415 BP; 136 A; 91 C; 80 G; 108 T; 0 other;  
  
Query Match 36.6%; Score 25.6; DB 22; Length 415;

Best Local Similarity 66.1%; Pred. No. 23;  
Matches 37; Conservative 0; Mismatches 19; Indels 0; Gaps 0;  
  
QY 15 AAATGGAACCTGAAATCAGATCCCTTCGCGGAATGGAACCTGAAATCAGATC 70  
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AC AAI04795;  
XX  
XX  
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XX  
XX Probe; human; breast disease; breast cancer; development disorder; ss;  
KW inflammatory disease; proliferative breast disease; non-carcinoma tumour.  
XX  
XX Homo sapiens.  
XX  
XX WO200157270-A2.  
XX  
XX  
PD 09-AUG-2001.  
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XX 29-JAN-2001; 2001WO-US00661.  
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XX 04-FEB-2000; 2000US-0180312.  
XX 26-MAY-2000; 2000US-0207456.  
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PA (MOLE-) MOLECULAR DYNAMICS INC.  
XX  
XX  
PI Penn SG, Hanzel DK, Chen W, Rank DR;  
XX WPI; 2001-476286/51.  
XX  
PT Novel single exon nucleic acid probe used to measuring gene expression  
in a human breast -  
XX  
XX  
PS Claim 25; SEQ ID No 4786; 322pp; English.  
XX  
XX The present invention relates to novel single exon nucleic acid probes.  
CC The present sequence is one such probe. The probes are useful for  
CC measuring human gene expression in a human breast sample, where the probe  
CC hybridises at high stringency to a nucleic acid expressed in the human  
CC breast. The probes are useful for predicting, diagnosing, grading,  
CC staging, monitoring and prognosing diseases of the human breast,  
CC particularly those diseases with polygenic aetiology. The diseases  
CC include: breast cancer, disorders of development, inflammatory diseases  
CC of the breast, fibrocystic changes, proliferative breast disease and  
CC non-carcinoma tumours.  
CC Note: The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences.  
XX  
XX  
SQ Sequence 415 BP; 136 A; 91 C; 80 G; 108 T; 0 other;  
  
Query Match 36.6%; Score 25.6; DB 22; Length 415;  
Best Local Similarity 66.1%; Pred. No. 23;  
Matches 37; Conservative 0; Mismatches 19; Indels 0; Gaps 0;  
  
QY 15 AAATGGAACCTGAAATCAGATCCCTTCGCGGAATGGAACCTGAAATCAGATC 70  
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RESULT 10  
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PR 08-DEC-2000; 2000US-0251868.  
PR 08-DEC-2000; 2000US-0251869.  
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PR 11-DEC-2000; 2000US-0254097.  
PR 05-JAN-2001; 2001US-0259678.  
XX  
PA (HUMA-) HUMAN GENOME SCI INC.  
XX  
PI Rosen CA, Barash SC, Ruben SM;  
XX WPI: 2001-502630/55.  
DR  
XX Polynucleotides encoding digestive system antigens, useful for  
PT diagnosing, treating, preventing and/or prognosing disorders of the  
PT digestive system, particularly cancer and cancer metastases -  
XX  
PS Disclosure; SEQ ID NO 4997; 986pp; English.  
XX The present invention provides the protein and coding sequences of a  
CC

PR 08-SEP-2000; 2000US-0231414.  
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PR 11-DEC-2000; 2000US-0254097.  
PR 05-JAN-2001; 2001US-0259678.  
XX (HUMA-) HUMAN GENOME SCI INC.  
XX Rosen CA, Barash SC, Ruben SM;  
XX WPI; 2001-502630/55.  
XX Polynucleotides encoding digestive system antigens, useful for  
XX diagnosing, treating, preventing and/or prognosing disorders of the  
XX digestive system, particularly cancer and cancer metastases -  
XX Disclosure; SEQ ID NO 4996; 986pp; English.  
XX The present invention provides the protein and coding sequences of a  
XX number of human digestive system antigens. These can be used in the  
XX diagnosis, treatment and prevention of digestive system disorders,  
XX including cancer, Meckel's diverticulum, bacterial or parasitic  
XX infections, appendicitis, Hirschsprung's disease, chronic colitis or  
XX ulcerative colitis. The present sequence is a genomic DNA fragment  
XX encoding a digestive system antigen of the invention.  
XX Sequence 551 BP; 187 A; 101 C; 116 G; 147 T; 0 other;  
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XX Query Match 38.6%; Score 27; DB 22; Length 551;  
XX Best Local Similarity 66.1%; Pred. No. 8.1;  
XX Matches 39; Conservative 0; Mismatches 20; Indels 0; Gaps 0;  
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ID AAK91421 standard; DNA; 551 BP.  
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XX 05-NOV-2001 (first entry)  
XX Human digestive system antigen genomic sequence SEQ ID NO: 4997.  
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XX Human; digestive system antigen; gene therapy; cancer; appendicitis;  
XX ulcerative colitis; infection; Hirschsprung's disease; chronic colitis;  
XX digestive system disorder; Meckel's diverticulum; ds.  
XX Homo sapiens.  
XX  
XX WO200155314-A2.  
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XX 02-AUG-2001.  
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XX 17-JAN-2001; 2001WO-US01324.  
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XX 31-JAN-2000; 2000US-0179065.  
XX 04-FEB-2000; 2000US-0180628.  
XX 24-FEB-2000; 2000US-0184664.  
XX 02-MAR-2000; 2000US-0186350.



CC (e.g. polyps and adenomas, intestinal inflammatory disorders, colitis,  
 CC colonic inflammation, diarrhea and dysentery, malabsorption syndromes,  
 CC (e.g. lactose intolerance), intestinal obstruction and sigmoid diseases.  
 CC The polynucleotide sequences of the invention can also be used in gene  
 CC therapy. AAS39582-AAS40060 represent DNA sequences encoding for the  
 CC novel human colon associated polypeptides of the invention.  
 CC Note: The sequence data for this patent did not form part of the printed  
 CC specification, but was obtained in electronic format directly from WIPO  
 CC at ftp.wipo.int/pub/published\_pct\_sequences.

XX Sequence 551 BP; 187 A; 101 C; 116 G; 147 T; 0 other;

SQ Query Match 38.6%; Score 27; DB 22; Length 551;

Best Local Similarity 66.1%; Pred. No. 8.1;

Matches 39; Conservative 0; Mismatches 20; Indels 0; Gaps 0;

QY 11 CGGGAATGGAATGGAATGAGATCCCTTCGCGGAATGGAATGGAATGAGAT 69

Db 193 CTGCTACTGAAACAAACAAATAGACCCCTGTTTGAGAAAGGAAATATGAAATCAGAT 251

# RESULT 4

40004

AAS40004 standard; DNA; 551 BP.

AC AAS40004;

XX DT 17-DEC-2001 (first entry)

DE Genomic sequence #423 encoding human colon associated polypeptide.

XX Human: colon cancer; congenital abnormality; infection; colitis;  
 KW inflammatory bowel disease; IBD; neoplastic disorder; gene therapy;  
 KW intestinal inflammatory disorder; malabsorption syndrome; gastric;  
 KW sigmoid disease; antibacterial; antiviral; antiinflammatory;  
 KW cytostatic; ds.

XX OS Homo sapiens.

XX PN WO200155302-A2.

XX PD 02-AUG-2001.

XX PF 17-JAN-2001; 2001WO-US01240.

XX PR 31-JAN-2000; 2000US-0179065.

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PR 19-MAY-2000; 2000US-0205515.

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PR 28-JUN-2000; 2000US-0214886.

PR 30-JUN-2000; 2000US-0215135.

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PR 05-JAN-2001; 2001US-0259678.  
XX  
XX (HUMA-) HUMAN GENOME SCI INC.  
XX  
XX Rosen CA, Barash SC, Ruben SM;  
XX  
XX WPI; 2001-465567/50.  
XX  
XX Isolated polypeptide for treating, preventing and/or prognosing  
XX disorders related to the colon including colon cancers and also for  
XX testing and detection e.g. diagnosis -  
XX  
XX Disclosure; SEQ ID No 900; 562pp; English.  
XX  
XX The present invention relates to the isolation of novel human colon  
XX associated polypeptides (AAU22468-AAU22701), and the cDNA and genomic  
XX sequences encoding for them. The sequences of the invention are useful  
XX in the diagnosis, treatment, prevention and/or prognosis of disorders  
XX of the colon including colon cancer, congenital abnormalities  
XX (e.g. atresia and stenosis), bacterial and viral infections,  
XX inflammatory bowel disease (IBD), neoplastic cell disorders



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OM nucleic - nucleic search, using sw model

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Listing first 45 summaries

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# SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
1	70	100.0	70	21	Firefly IRF-1 bind
2	27.6	39.4	2286	23	DNA encoding novel
3	27	38.6	551	22	Genomic sequence #
4	27	38.6	551	22	Genomic sequence #
5	27	38.6	551	22	Human digestive sy
6	27	38.6	551	22	Human digestive sy
c 7	26.2	37.4	481	24	Human ovarian canc
8	25.6	36.6	415	22	Human breast cell
9	25.6	36.6	415	22	Human foetal liver

10	25.6	36.6	415	22	ABA26396	Probe #4862 for ge
11	25.6	36.6	415	22	AAK04899	Human brain expres
12	25.6	36.6	415	22	AAK30423	Human bone marrow
13	25.6	36.6	415	22	AAI15029	Probe #4962 for ge
14	25.6	36.6	415	22	AAI16372	Probe #5058 used t
15	25.6	36.6	415	22	AAI04795	Probe #4786 used t
16	25.6	36.6	415	24	ABS05034	Human genome-deriv
17	25.6	36.6	8981	21	AAK69138	Human ABC1 gene ex
c 18	25.6	36.6	183999	22	AAF92831	Human ABC1 genomic
c 19	25.4	36.3	715	20	AAK37514	Human secreted pro
20	25.4	36.3	90104	23	ABL12402	Drosophila melanog
21	25.4	36.3	684707	24	ABQ87196	Listeria innocua c
22	25.4	36.3	3011208	24	ABQ69245	Listeria innocua c
c 23	25.2	36.0	348	22	AAK02035	Bladder cancer-ass
c 24	25.2	36.0	1230025	20	AAK91990	Nucleotide sequenc
25	25	35.7	976	23	ABV04483	Human prostate exp
26	24.8	35.4	310	20	AAV89187	EST clone CH699
c 27	24.8	35.4	685	21	AAK33345	Arabidopsis thalia
28	24.8	35.4	705	22	AAF22680	Human gastric canc
29	24.8	35.4	887	22	AAF22682	Human gastric canc
30	24.8	35.4	996	21	AAK05586	Streptococcus pneu
31	24.8	35.4	1789	20	AAK40059	Colon cancer assoc
32	24.8	35.4	2353	19	AAV40542	Homo sapiens secre
33	24.8	35.4	2566	22	AAF59641	Human cell cycle a
c 34	24.8	35.4	7568	23	ABL14768	Drosophila melanog
c 35	24.8	35.4	11887	19	AAV52279	Streptococcus pneu
c 36	24.8	35.4	49312	21	AAH51594	Human genomic sequ
37	24.8	35.4	140036	24	AAK98600	Human genomic DNA
c 38	24.6	35.1	1510	22	AAH99084	Human EST-derived
c 39	24.6	35.1	3884	22	AAH98627	Human EST-derived
c 40	24.6	35.1	5397	23	AAK56600	DNA encoding novel
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42	24.6	35.1	2365589	24	ABA90521	Genomic sequence o
43	24.4	34.9	244	22	AAK23933	Human ovarian PCR-
44	24.4	34.9	244	22	AAH82491	Human ovarian tumo
c 45	24.4	34.9	1555	21	AAK41543	Arabidopsis thalia

## ALIGNMENTS

### RESULT 1

AAK64592

ID AAC64592 standard; DNA; 70 BP.

AC AAC64592;

XX AAC64592;

XX AAC64592;

DT 15-FEB-2001 (first entry)

DE Firefly IRF-1 binding site nucleotide sequence SEQ ID NO:2.

XX Myeloproliferative sarcoma virus; MPSV; MPSV-E; enhancer; CMV;  
KW cytomagalovirus; firefly; IRF-1 binding site; minimal promoter;  
KW interferon regulatory factor 1 binding site; transcription regulator;  
KW promoter transactivator system; ds.  
XX Photinus pyralis.  
OS XX  
XX XX  
PN EPI046710-Al.  
XX XX  
PD 25-OCT-2000.  
XX XX  
XX 23-APR-1999; 99EP-0108068.  
XX XX  
PR 23-APR-1999; 99EP-0108068.  
XX XX  
PA (GBFB ) GBS BIOTECHNOLOGISCHE FORSCHUNG MBH.  
XX XX  
PI Mueller PP, Geserick C, Schroeder K, Hauser H;  
XX WPI; 2000-648930/63.  
DR WPI; 2000-648930/63.  
XX WPI; 2000-648930/63.  
PT Promoter-transactivator system, useful for inducing high level



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misc_feature /note="assembly_fragment"
2879. 6107
misc_feature /note="assembly_fragment"
6208. 7623
misc_feature /note="assembly_fragment"
clone_end:SP6
vector_side:right
7724. 11841
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clone_end:T7
vector_side:right
11942. 18456
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18557. 27382
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27483. 37124
misc_feature /note="assembly_fragment"
37225. 47276
misc_feature /note="assembly_fragment"
47377. 66713
misc_feature /note="assembly_fragment"
66814. 89140
misc_feature /note="assembly_fragment"
89241. 116360
misc_feature /note="assembly_fragment"
116461. 141816
misc_feature /note="assembly_fragment"
141917. 176775
misc_feature /note="assembly_fragment"
42683 a 45397 c 44545 g 42948 t 1202 others
ORIGIN
Query Match 41.78; Score 29.2; DB 2; Length 176775;
Best Local Similarity 74.08; Pred. No. 35;
Matches 37; Conservative 0; Mismatches 13; Indels 0; Gaps 0;

Qy 13 GGAATGGAACGAAATCAGACTCCCTCTCGGGAATGGAACCTGAAA 62
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 104829 GGAATGGAGACAGCACTGAGACTCTCCCTGTAAGGAAGAGAAA 104780

RESULT 15
AC067957 165373 bp DNA linear PRI 09-JAN-2002
LOCUS Homo sapiens BAC clone RP11-82K13 from 2, complete sequence.
DEFINITION AC067957
ACCESSION AC067957
VERSION AC067957.7 GI:15638744
KEYWORDS HTG.
SOURCE Homo sapiens.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Isak,A., Kozlowicz,A., Dixon,R. and Doeberer,A.
Sulston,J.E. and Waterston,R.
Toward a complete human genome sequence
Genome Res. 8 (11), 1097-1108 (1998)
95063792
PUBMED 9847074
REFERENCE 2 (bases 1 to 165373)
AUTHORS Isak,A., Kozlowicz,A., Dixon,R. and Doeberer,A.
TITLE The sequence of Homo sapiens BAC clone RP11-82K13
JOURNAL Unpublished (2001)
REFERENCE 3 (bases 1 to 165373)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (27-APR-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
REFERENCE 4 (bases 1 to 165373)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (18-SEP-2001) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,

```

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT

MO 63108, USA  
5 (bases 1 to 165373)  
Waterston,R.  
Direct Submission  
Submitted (09-JAN-2002) Department of Genetics, Washington  
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA  
On Sep 18, 2001 this sequence version replaced gi:14670148.  
----- Genome Center  
Center: Washington University Genome Sequencing Center  
Center code: WUGSC  
Web site: http://genome.wustl.edu/gsc  
Contact: sapiens@watson.wustl.edu  
----- Summary Statistics  
-----  
Center project name: H\_NH0082K13  
-----

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:  
all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

#### MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see http://genome.wustl.edu/gsc

#### SOURCE INFORMATION:

The RPCI-11 human BAC library was made from the blood of one male donor, as described by Osogawa,K., Woon,P.Y., Zhao,B., Frenken,E., Tatenos,M., Catanese,J.J. and de Jong,P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (http://www.resgen.com) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (http://bacpac.med.buffalo.edu)

VECTOR: pBACe3.6

#### NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is RP11-111K7; the clone sequenced to the right is RP11-89K21, 2000 bp overlap. Actual start of this clone is at base position 1 of RP11-82K13; actual end is at base position 29707 of RP11-89K21.

Data from AC012455, AC067791, and AC062030 was used to finish this clone, AC067957.

Polymorphisms exist between AC067957 and AC067791.

FEATURES  
source

Location/Qualifiers  
1. 165373  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/chromosome="2"  
/map="2"  
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/clone\_lib="RPCI-11"  
3097. 3427  
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misc\_feature

repeat\_region

repeat\_region

repeat\_region

misc\_feature

3590. 5549  
/note="match to EST AA485569 (NID:g2214788) ab09g06.rl"

<http://www.chori.org/bacpac/home.htm>  
 VECTOR: pBAC3.6

IMPORTANT: This sequence is not the entire insert of clone RP11-149P14. It may be shorter because we sequence overlapping sections only once, except for a short overlap.  
 The true left end of clone RP11-149P14 is at 1 in this sequence.  
 The true left end of clone RP1-539L13 is at 108122 in this sequence.  
 The true right end of clone RP11-192B7 is at 58622 in this sequence.

```

FEATURES             Location/Qualifiers
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                        /db_xref="taxon:9606"
                        /chromosome="1"
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                        /clone_lib="RPCI-11.1"
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                        /note="Sequence from overlapping clone RP11-192B7
                        (AL592314). Assembly confirmed by restriction digest."
     misc_feature      21038..21123
                        /note="Sequence from overlapping clone RP11-192B7
                        (AL592314). Assembly confirmed by restriction digest."
     misc_feature      50716..50718
                        /note="Sequence from overlapping clone RP11-192B7
                        (AL592314). Assembly confirmed by restriction digest."
     misc_feature      54763..54871
                        /note="Sequence from overlapping clone RP11-192B7
                        (AL592314). Assembly confirmed by restriction digest."
     misc_feature      86404..86414
                        /note="Sequence from uni-directional dGTP big dye
                        terminator reads only."
     misc_feature      86780..86903
                        /note="Single clone region. Reads generated from a
                        transposon library derived from a single pUC clone.
                        Restriction digest data confirm the assembly."
BASE COUNT            25978 a 28405 c 27853 g 27885 t
ORIGIN
Query Match          41.7%; Score 29.2; DB 9; Length 110121;
Best Local Similarity 74.0%; Pred. No. 35;
Matches 37; Conservative 0; Mismatches 13; Indels 0; Gaps 0;

Oy 13 GGAATGGAACTGAATCAGATCCCTCTCGGGAATCGAACTGAAA 62
||||| ||||| || ||||| ||||| ||||| ||||| |||||
Db 26583 GGAATGGAGACAGCAAGTCAGAACTCTCCCTGTAAAGGAAAGAGAAA 26534

RESULT 14
AC016978/c           176775 bp   DNA   linear   HTG 09-SEP-2000
LOCUS                Homo sapiens clone RP11-5N7, WORKING DRAFT SEQUENCE, 13 unordered
DEFINITION            pieces.
ACCESSION             AC016978
VERSION               AC016978.3 GI:10045327
KEYWORDS               HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE                Homo sapiens.
ORGANISM              Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE              1 (bases 1 to 176775)
AUTHORS               Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
                       Anderson,S., Baldwin,J., Barna,N., Beckerly,R., Bida,F.,
                       Boguslavskiy,L., Boukhgalter,B., Brown,A., Castile,A., Colangelo,M.,
                       Collins,S., Collymore,A., Cooke,P., DeArelano,K., Dewar,K.,
                       Domino,M., Doyle,M., Fenesor,J., Ferreira,P., FitzHugh,W.,
                       Forrest,C., Gage,D., Galagan,J., Gardyna,S., Grant,G., Hagos,B.,
                       Hearford,A., Horton,L., Howland,J.C., Johnson,R., Jones,C., Kann,L.,
                       Karatsas,A., Klein,J., Landers,T., Lehoczy,J., Lieu,C., Locke,K.,
                       Macdonald,P., Marquis,N., McEwan,P., McGurk,A., McKernan,K.,
                       2 (bases 1 to 176775)
AUTHORS               Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
                       Anderson,S., Baldwin,J., Barna,N., Beckerly,R., Bida,F.,
                       Boguslavskiy,L., Boukhgalter,B., Brown,A., Castile,A., Colangelo,M.,
                       Collins,S., Collymore,A., Cooke,P., DeArelano,K., Dewar,K.,
                       Domino,M., Doyle,M., Fenesor,J., Ferreira,P., FitzHugh,W.,
                       Forrest,C., Gage,D., Galagan,J., Gardyna,S., Grant,G., Hagos,B.,
                       Hearford,A., Horton,L., Howland,J.C., Johnson,R., Jones,C., Kann,L.,
                       Karatsas,A., Klein,J., Landers,T., Lehoczy,J., Lieu,C., Locke,K.,
                       Macdonald,P., Marquis,N., McEwan,P., McGurk,A., McKernan,K.,

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Meldrim,J., Morrow,J., Naylor,J., Norman,C.H., O'Connor,T.,
O'Donnell,P., Peterson,K., Pierre,N., Pollara,V., Riley,R.,
Rothman,D., Roy,A., Santos,R., Severy,P., Stange-Thomann,N.,
Stojanovic,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
Tirrell,A., Vassiliev,H., Viel,R., VO.A., Wu,X., Wyman,D., Ye.W.J.,
Zimmer,A. and Zody,M.
Direct Submission
Submitted (09-DEC-1999) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Sep 9, 2000 this sequence version replaced gi:6648261.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L2875
Center clone name: 5_N7
----- Summary Statistics
Sequencing vector: M13; M77815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 166773 bases at least Q40
Consensus quality: 172242 bases at least Q30
Consensus quality: 174181 bases at least Q20
Insert size: 85000; agarose-fp
Insert size: 175575; sum-of-contigs
Quality coverage: 8.8 in Q20 bases; agarose-fp
Quality coverage: 4.3 in Q20 bases; sum-of-contigs
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* NOTE: This is a 'working draft' sequence. It currently
* consists of 13 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
* 1 2778: contig of 2778 bp in length
* 2779 2878: gap of 100 bp
* 2879 6107: contig of 3229 bp in length
* 6108 6207: gap of 100 bp
* 6208 7623: contig of 1416 bp in length
* 7624 7723: gap of 100 bp
* 7724 11841: contig of 4118 bp in length
* 11842 11941: gap of 100 bp
* 11942 18456: contig of 6515 bp in length
* 18457 18556: gap of 100 bp
* 18557 27382: contig of 8826 bp in length
* 27383 27482: gap of 100 bp
* 27483 37124: contig of 9642 bp in length
* 37125 37224: gap of 100 bp
* 37225 47276: contig of 10052 bp in length
* 47277 47376: gap of 100 bp
* 47377 66713: contig of 19337 bp in length;
* 66714 66813: gap of 100 bp
* 66814 89140: contig of 22327 bp in length
* 89141 89240: gap of 100 bp
* 89241 116360: contig of 27120 bp in length
* 116361 116460: gap of 100 bp
* 116461 141816: contig of 25356 bp in length
* 141817 141916: gap of 100 bp
* 141917 176775: contig of 34859 bp in length.
Location/Qualifiers
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/db_xref="taxon:9606"
/clone="RP11-5N7"
/clone_lib="RPCI-11 Human Male BAC"
misc_feature 1..2778

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\* 25929 26799: contig of 871 bp in length  
 \* 26800 26899: gap of 100 bp  
 \* 26900 27808: contig of 909 bp in length  
 \* 27809 27908: gap of 100 bp  
 \* 27909 28812: contig of 904 bp in length  
 \* 28813 28912: gap of 100 bp  
 \* 28913 29813: contig of 901 bp in length  
 \* 29814 29913: gap of 100 bp  
 \* 29914 30807: contig of 894 bp in length  
 \* 30808 30907: gap of 100 bp  
 \* 30908 31790: contig of 883 bp in length  
 \* 31791 31890: gap of 100 bp  
 \* 31891 32777: contig of 887 bp in length  
 \* 32778 32877: gap of 100 bp  
 \* 32878 33789: contig of 912 bp in length  
 \* 33790 33889: gap of 100 bp  
 \* 33890 34792: contig of 903 bp in length  
 \* 34793 34892: gap of 100 bp  
 \* 34893 35795: contig of 903 bp in length  
 \* 35796 35895: gap of 100 bp  
 \* 35896 36798: contig of 903 bp in length  
 \* 36799 36898: gap of 100 bp  
 \* 36899 37791: contig of 893 bp in length  
 \* 37792 37891: gap of 100 bp  
 \* 37892 38732: contig of 841 bp in length  
 \* 38733 38832: gap of 100 bp  
 \* 38833 39745: contig of 914 bp in length  
 \* 39747 39846: gap of 100 bp  
 \* 39847 40745: contig of 899 bp in length  
 \* 40746 40845: gap of 100 bp  
 \* 40846 41743: contig of 898 bp in length  
 \* 41744 41843: gap of 100 bp  
 \* 41844 42723: contig of 880 bp in length  
 \* 42724 42823: gap of 100 bp  
 \* 42824 43687: contig of 864 bp in length  
 \* 43688 43787: gap of 100 bp  
 \* 43788 44683: contig of 896 bp in length  
 \* 44684 44783: gap of 100 bp  
 \* 44784 45688: contig of 902 bp in length  
 \* 45689 45785: gap of 100 bp  
 \* 45786 46689: contig of 904 bp in length  
 \* 46690 46789: gap of 100 bp  
 \* 46790 47690: contig of 901 bp in length  
 \* 47691 47790: gap of 100 bp  
 \* 47791 48690: contig of 900 bp in length  
 \* 48691 48790: gap of 100 bp  
 \* 48791 49709: contig of 919 bp in length  
 \* 49710 49809: gap of 100 bp  
 \* 49810 50684: contig of 875 bp in length  
 \* 50685 50784: gap of 100 bp  
 \* 50785 51684: contig of 900 bp in length  
 \* 51685 51784: gap of 100 bp  
 \* 51785 52685: contig of 901 bp in length  
 \* 52686 52785: gap of 100 bp  
 \* 52786 53691: contig of 906 bp in length  
 \* 53692 53791: gap of 100 bp  
 \* 53792 54688: contig of 897 bp in length  
 \* 54689 54788: gap of 100 bp  
 \* 54789 55703: contig of 917 bp in length  
 \* 55706 55805: gap of 100 bp  
 \* 55806 56714: contig of 909 bp in length  
 \* 56715 56814: gap of 100 bp  
 \* 56815 57704: contig of 890 bp in length  
 \* 57705 57804: gap of 100 bp  
 \* 57805 58702: contig of 898 bp in length  
 \* 58703 58802: gap of 100 bp  
 \* 58803 59683: contig of 881 bp in length  
 \* 59684 59783: gap of 100 bp  
 \* 59784 60662: contig of 879 bp in length  
 \* 60663 60762: gap of 100 bp  
 \* 60763 61642: contig of 880 bp in length  
 \* 61643 61742: gap of 100 bp  
 \* 61743 62617: contig of 875 bp in length

\* 62618 62717: gap of 100 bp  
 \* 62718 63623: contig of 906 bp in length  
 \* 63624 63723: gap of 100 bp  
 \* 63724 64617: contig of 894 bp in length  
 \* 64618 64717: gap of 100 bp  
 \* 64718 65635: contig of 918 bp in length  
 \* 65636 65735: gap of 100 bp  
 \* 65736 66612: contig of 877 bp in length  
 \* 66613 66712: gap of 100 bp  
 \* 66713 67591: contig of 879 bp in length  
 \* 67592 67691: gap of 100 bp  
 \* 67692 68598: contig of 907 bp in length  
 \* 68599 68698: gap of 100 bp  
 \* 68699 69594: contig of 896 bp in length  
 \* 69595 69694: gap of 100 bp  
 \* 69695 70501: contig of 807 bp in length  
 \* 70502 70601: gap of 100 bp  
 \* 70602 71452: contig of 851 bp in length  
 \* 71453 71552: gap of 100 bp  
 \* 71553 72370: contig of 818 bp in length

Query Match 42.3%; Score 29.6; DB 2; Length 90747;  
 Best Local Similarity 88.9%; Pred. NO. 26;  
 Matches 32; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

OY 4 CCCTTCTCGGGAATGGAACCTGAAATCAGATCCC 39  
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 Db 1466 CCCTTCTCGGGAATGGAACCTGAAATCAGATCCC 1431

RESULT 13  
 AL590989/LOCUS  
 DEFINITION Human DNA sequence from clone RP11-149P14 on chromosome 1, complete sequence.  
 ACCESSION AL590989  
 VERSION AL590989.15 GI:16973092  
 KEYWORDS HTG  
 SOURCE human.  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 REFERENCE 1 (bases 1 to 110121)  
 AUTHORS Howden, P.  
 TITLE Direct Submission  
 JOURNAL Submitted (15-NOV-2001) Wellcome Trust Sanger Institute, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk  
 COMMENT On Nov 16, 2001 this sequence version replaced gi:16508264.  
 During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.  
 This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em, EMBL; Sw, SWISSPROT; Tr, TREMBL; Wp, WORMPEP; Information on the WORMPEP database can be found at  
[http://www.sanger.ac.uk/Projects/C\\_elegans/wormpep](http://www.sanger.ac.uk/Projects/C_elegans/wormpep)  
 This sequence was generated from part of bacterial clone contigs of human chromosome 1, constructed by the Sanger Centre Chromosome 1 Mapping Group. Further information can be found at  
<http://www.sanger.ac.uk/HGP/Chrl>  
 RP11-149P14 is from the library RPCI-11.1 constructed by the group of Pieter de Jong. For further details see





Center: Whitehead Institute/ MIT Center for Genome Research  
Center code: WIBR  
Web site: <http://www-seq.wi.mit.edu>  
Contact: [sequence\\_submissions@genome.wi.mit.edu](mailto:sequence_submissions@genome.wi.mit.edu)  
----- Project Information  
Center project name: L5392  
Center clone name: 127 H 5

## FEATURES

## source

JURES	source	Location/Qualifiers
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		/db_xref="taxon:9606"
		/chromosome="8"
		/map="8"
		/clone="RP11-127H5"
		/cvalue_lib="RPC1-II Human Male BAC"
repeat_region		2879..3178
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repeat_region		3814..3842
repeat_region		/rpt_family="AT_Rich"
repeat_region		complement(4030..4462)
repeat_region		/rpt_family="L2"
repeat_region		4486..4540
repeat_region		/rpt_family="(TTCA)n"
repeat_region		complement(4550..4830)
repeat_region		/rpt_family="L1Mc1"
repeat_region		4831..4920
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repeat_region		/rpt_family="L1Mc1"
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repeat_region		/rpt_family="MIR3"
repeat_region		6901..6927
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repeat_region		14722..14907
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repeat_region		14923..15077
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repeat_region		complement(18749..18863)

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repeat_region	/rpt_family="MIR" complement(20999..21095)
repeat_region	/rpt_family="L2" 21284..21520
repeat_region	/rpt_family="MLT1J1" 21949..21991
repeat_region	/rpt_family="(TTTTG)n" 23647..23713
repeat_region	/rpt_family="(CA)n" 24203..24340
repeat_region	/rpt_family="MIR" complement(24837..25022)
repeat_region	/rpt_family="MIR3" complement(25220..25371)
repeat_region	/rpt_family="L2" 25505..25811
repeat_region	/rpt_family="ERVLF-F" 25815..26206
repeat_region	/rpt_family="MLT2F" 26278..26373
repeat_region	/rpt_family="MLT2F" 26395..26663
repeat_region	/rpt_family="L1PAL6" 26664..26953
repeat_region	/rpt_family="AluSx" 26954..27438

Query Match 42.9%; Score 30; DB 9; Length 150883;  
Best Local Similarity 72.2%; Pred. No. 19;  
Matches 39; Conservative 0; Mismatches 15; Indels: 0

[illegible]

RESULT 11

AL591711	197743 bp	DNA	linear	ROD 25-MAY-2002					
LOCUS	Mouse DNA sequence from clone RP23-19L12 on chromosome 2, complete sequence.								
ACCESSION	AL591711								
VERSION	1.22	GI:21217770							
KEYWORDS	HTG.								
SOURCE	house mouse.								
ORGANISM	Mus musculus								
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.								
AUTHORS	1 (bases 1 to 197743) Johnson, C.								
TITLE	Direct Submission								
JOURNAL	Submitted (25-MAY-2002) Wellcome Trust Sanger Institute, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk								
COMMENT	On May 27, 2002 this sequence version replaced gi:21068504. During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above. This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate								





\* 171141 207659: contig of 36519 bp in length  
 \* 207660 207759: gap of 100 bp  
 \* 207760 270196: contig of 62437 bp in length  
 \* 270197 270296: gap of 100 bp  
 \* 270297 270795: contig of 499 bp in length.

Query Match	43.7%	Score	30.6	DB 2	Length	270795
Best Local Similarity	80.0%	Pred. No.	13			
Matches	36	Conservative	0	Mismatches	9	Indels
					0	Gaps
					0	

RESULT	8
AC022602/c	
LOCUS	62795 bp DNA linear HTG 12-APR-2001
DEFINITION	Homo sapiens chromosome 8 clone RP11-700J23 map 8, LOW-PASS SEQUENCE SAMPLING.

Anderson S., Baldwin J., Barna N., Beckerly R., Beda F., Boguslavsky L., Boukhalter B., Brown A., Burkett G., Castle A., Chospel Y., Collangelo M., Collins S., Collymore A., Cooke P., DeArrellano K., Dewar K., Domino M., Doyle M., Fenestor J., Ferreira P., FitzHugh W., Forrest C., Gage D., Galagan J., Gardyna S., Grant G., Hagos B., Heaford A., Horton L., Howland J.C., Johnson R., Jones C., Kann L., Karatas A., Klein J., Landers T., Lehoczyk J., Levine R., Lieu C., Liu G., Locke K., Macdonald P., Marquis N., McEwan P., McGurk A., McKernan K., McPheeters R., Meldrum J., Meneus B., Morrow J., Naylor J., Norman C.H., O'Connor T., O'Donnell P., Oliver T.M., Peterson K., Pierre N., Pisani C., Pollara V., Raymond C., Riley R., Rothman D., Roy A., Santos R., Severy P., Spencer B., Stange-Thomann N., Stojanovic N., Subramanian A., Talamas J., Tesfaye S., Theodore J., Tirtrell A., Vassiliev H., Viel R., Vo A., Wu X., Wyman D., Ye W.J., Zimmer A. and Zody M.

Direct Submission  
Submitted (06-FEB-2000) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA.  
On Apr 12, 2001 this sequence version replaced gi:11990712.  
All repeats were identified using RepeatMasker:

\* NOTE: This record contains 73 individual  
\* sequencing reads that have not been assembled into  
\* contigs. Runs of N are used to separate the reads  
\* and the order in which they appear is completely  
\* arbitrary. Low-pass sequence sampling is useful for  
\* identifying clones that may be gene-rich and allows  
\* overlap relationships among clones to be deduced.  
\* However, it should not be assumed that this clone  
\* will be sequenced to completion. In the event that  
\* the record is updated, the accession number will  
\* be preserved.

TITLE	JOURNAL	COMMENT
-------	---------	---------

Direct Submission  
Submitted (06-FEB-2000) Whitehead Institute/MIT Center for Genomics  
Research, 320 Charles Street, Cambridge, MA 02141, USA.  
On April 12, 2001 this sequence version replaced gi:11950712.  
All repeats were identified using RepeatMasker:  
Smit, A.F.A. & Green, P. (1996-1397)  
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>  
----- Genome Center



SOURCE Norway rat.  
ORGANISM Rattus norvegicus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;  
Rattus.

REFERENCE 1 (bases 1 to 196360)  
AUTHORS Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-osman,F.R., Allen,C.,  
Alsbrooks,S.L., Amarantunge,H.C., Are,J.R., Ayele,M., Banks,T.,  
Barbosa,J., Benton,J., Bimaga,K., Blankenburg,K., Bonnin,D.,  
Bouck,J., Bowie,S., Brileva,M., Brown,E., Brown,M., Bryant,N.P.,  
Buhay,C., Burch,P., Burkett,C., Burrell,K.L., Byrd,N.C.,  
Carroll,T.F., Carter,M., Cavazos,S.R., Chacko,J., Chavez,D.,  
Chen,G., Chen,R., Chen,Z., Chowdhry,I., Christopoulos,C.,  
Cleveland,C.D., Cox,C., Coyle,M.D., Dathorne,S.R., David,R.,  
Devila,M.L., Davis,C., Davy-Carroll,L., Dederich,D.A.,  
DeLaney,K.R., Delgado,O., Denn,A.L., Ding,Y., Dinh,H.H.,  
Douthwaite,K.J., Draper,H., Dugan-Rocha,S., Durbin,K.J.,  
Earnhart,C., Edgar,D., Edwards,C.C., Elhaj,C., Escotto,M.,  
Falls,T., Ferraguto,D., Flagg,N., Ford,J., Foster,P., Frantz,P.,  
Gabisi,A., Gao,J., Garcia,A., Garner,T., Garza,N., Gill,R.,  
Gorrell,J.H., Guevara,W., Gunaratne,P., Hale,S., Hamilton,K.,  
Harris,C., Harris,K., Hart,M., Havlak,P., Hawes,A., Hernandez,J.,  
Hernandez,O., Hodgson,A., Hoques,M., Holloway,C., Hollins,B.,  
Honsi,F., Howard,S., Huber,J., Hulyk,S., Hume,J., Jackson,L.E.,  
Jacobson,B., Jia,Y., Johnson,R., Jolivet,S., Joudah,S.,  
Karlisson,E., Kelly,S., Khan,U., King,L., Korvan,J., Kovar,C.,  
Kratovic,J., Kureshi,A., Landry,N., Leal,B., Lewis,L.C., Lewis,L.,  
Li,J., Li,Z., Lichtarge,O., Lieu,C., Liu,J., Liu,W., Louissegh,H.,  
Lozado,R.J., Lu,X., Lucier,A., Lucier,R., Luna,R., Ma,J.,  
Maheshwari,M., Mapua,P., Martin,R., Martindale,A., Martinez,E.,  
Massey,E., Mawhney,E., McLeod,M.P., Meador,M., Mei,G., Metzker,M.,  
Miner,G., Miner,Z., Mitchell,T., Mohabbat,K., Morgan,M., Morris,S.,  
Moser,M., Neal,D., Newton,J., Newton,N., Nguyen,A., Nguyen,N.,  
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Oraguene,N., Oviedo,R., Pace,A., Payton,B., Peery,J., Perez,L.,  
Peters,L., Pickens,R., Primus,E., Pu,L.L., Quiles,M., Ren,Y.,  
Rives,M., Rojas,A., Rojibokan,I., Rolfe,M., Ruiz,S., Savery,G.,  
Scherer,S., Scott,G., Shen,H., Shoostari,N., Sisson,I.,  
Sodergren,E., Sonake,T., Sparks,A., Stanley,H., Stone,H.,  
Sutton,A., Svatek,A., Tabor,P., Tamerisa,A., Tamerisa,K., Tang,H.,  
Tansey,J., Taylor,C., Taylor,T., Telford,B., Thomas,N., Thomas,S.,  
Usmani,K., Vasquez,L., Vera,V., Villalon,D., Vinson,R., Wang,Q.,  
Wang,S., Ward-Moore,S., Warren,R., Washington,C., Watlington,S.,  
Williams,G., Williamson,A., Wleczyk,R., Wooden,S., Worley,K.,  
Wu,C., Wu,Y., Wu,Y.F., Zhou,J., Zorrilla,S., Nelson,D.,  
Weinstock,G. and Gibbs,R.

Direct Submission  
Unpublished  
2 (bases 1 to 196360)  
Worley,K.C.  
Direct Submission  
Submitted (24-OCT-2001) Human Genome Sequencing Center, Department  
of Molecular and Human Genetics, Baylor College of Medicine, One  
Baylor Plaza, Houston, TX 77030, USA

REFERENCE 3 (bases 1 to 196360)  
AUTHORS Worley,K.C.  
TITLE Direct Submission  
JOURNAL Submitted (12-JUL-2002) Human Genome Sequencing Center, Department  
of Molecular and Human Genetics, Baylor College of Medicine, One  
Baylor Plaza, Houston, TX 77030, USA

COMMENT On Jul 11, 2002 this sequence version replaced gi:17973146.  
----- Genome Center  
Center: Baylor College of Medicine  
Center code: BCM  
Web site: <http://www.hgsc.bcm.tmc.edu/>  
Contact: [hgsc-help@bcm.tmc.edu](mailto:hgsc-help@bcm.tmc.edu)  
----- Project Information  
Center project name: GHFL  
Center clone name: CH230-64F11  
----- Summary Statistics  
Sequencing vector: Plasmid;  
Chemistry: dye-terminator Big Dye; 100% of reads  
Assembly program: Phrap; version 0.990329

Consensus quality: 130985 bases at least Q40  
Consensus quality: 137495 bases at least Q30  
Consensus quality: 142650 bases at least Q20  
-----  
\* NOTE: Estimated insert size may differ from sequence length  
(see [http://www.hgsc.bcm.tmc.edu/docs/Genbank\\_draft\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)).  
\* NOTE: This is a 'working draft' sequence. It currently  
\* consists of 78 contigs. The true order of the pieces  
\* is not known and their order in this sequence record is  
\* arbitrary. Gaps between the contigs are represented as  
\* runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence  
\* as soon as it is available and the accession number will  
\* be preserved.

1  
\* 1005: contig of 1005 bp in length  
\* 1105: gap of unknown length  
\* 1106: contig of 1015 bp in length  
\* 2121: gap of unknown length  
\* 2221: contig of 1051 bp in length  
\* 3272: gap of unknown length  
\* 3372: contig of 1571 bp in length  
\* 4943: gap of unknown length  
\* 5042: contig of 1653 bp in length  
\* 6696: gap of unknown length  
\* 6795: contig of 1687 bp in length  
\* 8483: gap of unknown length  
\* 8583: contig of 1029 bp in length  
\* 9612: gap of unknown length  
\* 9712: contig of 1091 bp in length  
\* 10802: contig of 1091 bp in length  
\* 10902: gap of unknown length  
\* 11944: contig of 1042 bp in length  
\* 12045: gap of unknown length  
\* 13423: contig of 1379 bp in length  
\* 13523: gap of unknown length  
\* 14596: contig of 1073 bp in length  
\* 14597: gap of unknown length  
\* 16601: contig of 1905 bp in length  
\* 16701: gap of unknown length  
\* 17907: contig of 1206 bp in length  
\* 18007: gap of unknown length  
\* 18008: contig of 1605 bp in length  
\* 19614: gap of unknown length  
\* 19713: contig of 1113 bp in length  
\* 20826: gap of unknown length  
\* 20927: contig of 1632 bp in length  
\* 22558: gap of unknown length  
\* 24413: contig of 1755 bp in length  
\* 24513: gap of unknown length  
\* 25973: contig of 1460 bp in length  
\* 26073: gap of unknown length  
\* 27357: contig of 1284 bp in length  
\* 27458: gap of unknown length  
\* 28672: contig of 1215 bp in length  
\* 28673: gap of unknown length  
\* 30074: contig of 1302 bp in length  
\* 30174: gap of unknown length  
\* 31717: contig of 1543 bp in length  
\* 31718: gap of unknown length  
\* 31818: contig of 1361 bp in length  
\* 33279: gap of unknown length  
\* 34465: contig of 1187 bp in length  
\* 34565: gap of unknown length  
\* 37040: contig of 2475 bp in length  
\* 37140: gap of unknown length  
\* 38197: contig of 1057 bp in length  
\* 38297: gap of unknown length  
\* 40231: contig of 1934 bp in length  
\* 40331: gap of unknown length  
\* 42055: contig of 1724 bp in length  
\* 42056: gap of unknown length  
\* 43274: contig of 1119 bp in length  
\* 43374: gap of unknown length  
\* 45875: contig of 2501 bp in length







Insert size: 158298; sum-of-contigs

Quality coverage.

\* NOTE: This is a 'working draft' sequence. It currently  
\* consists of 30 contigs. The true order of the pieces  
\* is not known and their order in this sequence record is  
\* arbitrary. Gaps between the contigs are represented as  
\* runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence  
\* as soon as it is available and the accession number will  
\* be preserved.

\* 1 1115: contig of 1115 bp in length  
\* 1116 1215: gap of 100 bp  
\* 1216 2414: contig of 1199 bp in length  
\* 2415 2514: gap of 100 bp  
\* 2515 3561: contig of 1047 bp in length  
\* 3562 3661: gap of 100 bp  
\* 3662 4929: contig of 1268 bp in length  
\* 4930 5029: gap of 100 bp  
\* 5030 6039: contig of 1010 bp in length  
\* 6040 6139: gap of 100 bp  
\* 6140 7469: contig of 1330 bp in length  
\* 7470 7569: gap of 100 bp  
\* 7570 9736: contig of 2167 bp in length  
\* 9737 9836: gap of 100 bp  
\* 9837 12165: contig of 2329 bp in length  
\* 12166 12265: gap of 100 bp  
\* 12266 12623: contig of 358 bp in length  
\* 12624 12723: gap of 100 bp  
\* 12724 14465: contig of 1742 bp in length  
\* 14466 14565: gap of 100 bp  
\* 14566 35963: contig of 1398 bp in length  
\* 15964 16063: gap of 100 bp  
\* 16064 18074: contig of 2011 bp in length  
\* 18075 18174: gap of 100 bp  
\* 18175 21150: contig of 2976 bp in length  
\* 21151 21250: gap of 100 bp  
\* 21251 22852: contig of 1602 bp in length  
\* 22853 22952: gap of 100 bp  
\* 22953 25375: contig of 2423 bp in length  
\* 25376 25475: gap of 100 bp  
\* 25476 27959: contig of 2484 bp in length  
\* 27960 28059: gap of 100 bp  
\* 28060 30399: contig of 2340 bp in length  
\* 30400 30499: gap of 100 bp  
\* 30500 34315: contig of 3816 bp in length  
\* 34316 34415: gap of 100 bp  
\* 34416 37936: contig of 3521 bp in length  
\* 37937 38036: gap of 100 bp  
\* 38037 40122: contig of 2086 bp in length  
\* 40123 40222: gap of 100 bp  
\* 40223 43895: contig of 3673 bp in length  
\* 43896 43995: gap of 100 bp  
\* 43996 49451: contig of 5456 bp in length  
\* 49452 49551: gap of 100 bp  
\* 49552 54107: contig of 4556 bp in length  
\* 54108 54207: gap of 100 bp  
\* 54208 60101: contig of 5894 bp in length  
\* 60102 60201: gap of 100 bp  
\* 60202 64961: contig of 4760 bp in length  
\* 64962 65061: gap of 100 bp  
\* 65062 75980: contig of 10919 bp in length  
\* 75981 76080: gap of 100 bp  
\* 76081 89332: contig of 13252 bp in length  
\* 89333 89432: gap of 100 bp  
\* 89433 104747: contig of 15315 bp in length  
\* 104748 104847: gap of 100 bp  
\* 104848 120535: contig of 15708 bp in length  
\* 120536 120556: gap of 100 bp  
\* 120556 161198: contig of 40543 bp in length.

Location/Qualifiers

1. .161198  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"

FEATURES  
source

misc\_feature /clone="RP11-71A1"  
1. .1115 /clone\_lib="RPC1-11 Human Male BAC"  
misc\_feature /note="assembly\_fragment"  
1216. 2414  
misc\_feature /note="assembly\_fragment"  
2515. .3561  
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3662. .4929  
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5030. 6039  
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6140. .7469  
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7570. .9736  
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9837. 12165  
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vector\_side:right  
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28060. .30399  
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54208. .60101  
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60202. .64961  
misc\_feature /note="assembly\_fragment"  
65062. .75980  
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misc\_feature /note="assembly\_fragment"  
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104848. .120555  
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120556. .161198  
misc\_feature /note="assembly\_fragment"  
2901 others  
BASE COUNT 40700 a 38352 c 37941 g 41304 t  
ORIGIN

Query Match 45.1%; Score 31.6; DB 2; Length 161198;  
Best Local Similarity 74.1%; Pred. No. 5.9;  
Matches 40; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

\* 27254 28110: contig of 857 bp in length  
\* 28111 28210: gap of 100 bp  
\* 28211 29057: contig of 847 bp in length  
\* 29058 29157: gap of 100 bp  
\* 29158 30015: contig of 858 bp in length  
\* 30016 30115: gap of 100 bp  
\* 30116 30959: contig of 844 bp in length  
\* 30960 31059: gap of 100 bp  
\* 31060 31892: contig of 833 bp in length  
\* 31893 31992: gap of 100 bp  
\* 31993 32825: contig of 833 bp in length  
\* 32826 32925: gap of 100 bp  
\* 32926 33786: contig of 861 bp in length  
\* 33787 33886: gap of 100 bp  
\* 33887 34730: contig of 844 bp in length  
\* 34731 34830: gap of 100 bp  
\* 34831 35686: contig of 856 bp in length  
\* 35687 35788: gap of 100 bp  
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\* 35843 36742: gap of 100 bp  
\* 36743 37588: contig of 846 bp in length  
\* 37589 37688: gap of 100 bp  
\* 37689 38583: contig of 895 bp in length  
\* 38584 38683: gap of 100 bp  
\* 38684 39538: contig of 855 bp in length  
\* 39539 39638: gap of 100 bp  
\* 39639 40465: contig of 827 bp in length  
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\* 40566 41398: contig of 833 bp in length  
\* 41399 41498: gap of 100 bp  
\* 41499 42350: contig of 852 bp in length  
\* 42351 42450: gap of 100 bp  
\* 42451 43323: contig of 873 bp in length  
\* 43324 43423: gap of 100 bp  
\* 43424 44274: contig of 851 bp in length  
\* 44275 44374: gap of 100 bp  
\* 44375 45243: contig of 869 bp in length  
\* 45244 45343: gap of 100 bp  
\* 45344 46198: contig of 855 bp in length  
\* 46199 46298: gap of 100 bp  
\* 46299 47210: contig of 912 bp in length  
\* 47211 47310: gap of 100 bp  
\* 47311 48149: contig of 839 bp in length  
\* 48150 48249: gap of 100 bp  
\* 48250 49095: contig of 846 bp in length  
\* 49096 49195: gap of 100 bp  
\* 49196 50025: contig of 830 bp in length  
\* 50026 50128: gap of 100 bp  
\* 50128 50955: contig of 830 bp in length  
\* 50956 51055: gap of 100 bp  
\* 51056 51889: contig of 834 bp in length  
\* 51890 51989: gap of 100 bp  
\* 51990 52860: contig of 871 bp in length  
\* 52861 52960: gap of 100 bp  
\* 52961 53793: contig of 833 bp in length  
\* 53794 53893: gap of 100 bp  
\* 53894 54738: contig of 845 bp in length  
\* 54739 54838: gap of 100 bp  
\* 54839 55682: contig of 844 bp in length  
\* 55683 55782: gap of 100 bp  
\* 55783 55640: contig of 858 bp in length  
\* 55641 56740: gap of 100 bp  
\* 56741 57576: contig of 836 bp in length  
\* 57577 57676: gap of 100 bp  
\* 57677 58527: contig of 851 bp in length  
\* 58528 58627: gap of 100 bp  
\* 58628 59468: contig of 841 bp in length  
\* 59469 59568: gap of 100 bp  
\* 59569 60400: contig of 832 bp in length  
\* 60401 60500: gap of 100 bp  
\* 60501 61390: contig of 890 bp in length  
\* 61391 61490: gap of 100 bp  
\* 61491 62346: contig of 856 bp in length

\* 62347 62446: gap of 100 bp  
\* 62447 63310: contig of 864 bp in length  
\* 63311 63410: gap of 100 bp  
\* 63411 64262: contig of 852 bp in length  
\* 64263 64362: gap of 100 bp  
\* 64363 65192: contig of 830 bp in length  
\* 65193 65292: gap of 100 bp

Query Match 45.1%; Score 31.6; DB 2; Length 69861;  
Best Local Similarity 74.1%; Pred. No. 5.7;  
Matches 40; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

QY 14 GAATGGAACAGAAATCAGATCCCTTCGGGAATGGAACAGAAATCAG 67  
||||| ||||||| ||| ||||||| ||| ||||||| ||||||| ||| |||  
DB 1327 GAATGGAACAGAAATCAGATCCCTTCGGGAATGGAATCTTGAATCTTAG 1380

RESULT 3  
AC015867  
LOCUS Homo sapiens clone RP11-71A1, WORKING DRAFT SEQUENCE, 30 unordered  
DEFINITION pieces.  
ACCESSION AC015867  
VERSION AC015867.2 GI:7249301  
KEYWORDS HTG; HTGS\_PHASE1; HTGS\_DRAFT.  
SOURCE Homo sapiens  
ORGANISM Homo sapiens

REFERENCE  
AUTHORS Birren,B., Linton,L., Nusbaum,C. and Lander,E.  
TITLE 1 (bases 1 to 161198)  
JOURNAL Homo sapiens, clone RP11-71A1  
REFERENCE 2 (bases 1 to 161198)  
AUTHORS Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,  
Baldwin,J., Barna,N., Beckerly,R., Boquslavkiy,L., Boukhgalter,B.,  
Brown,A., Castle,A., Colangelo,M., Collins,S., Collymore,A., Cooke,P.,  
Dearellano,K., Dewar,K., Domino,M., Donelan,L., Doyle,M., Ferreira,P.,  
FitzHugh,W., Forrest,C., Funke,R., Gage,D., Galagan,J., Gardyna,S.,  
Grant,G., Hago,B., Heaford,A., Horton,L., Howland,J.C., Johnson,R.,  
Jones,C., Kann,L., Karatas,A., Klein,J., Lehoczy,J., Lieu,C., Locke,K.,  
Macdonald,P., Marquis,N., McEwan,P., McGurk,A., McKernan,K.,  
McLaughlin,J., Meldrum,J., Morrow,J., Naylor,J., Norman,C.H.,  
O'Connor,T., O'Donnell,P., Peterson,K., Pollara,V., Riley,R., Roy,A., Santos,R., Severy,P.,  
Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J., Testaye,S.,  
Tirrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X., Wyman,D., Ye,W.J.,  
Zimmer,A. and Zody,M.

Direct Submission  
Submitted (17-NOV-1999) Whitehead Institute/MIT Center for Genome  
Research, 320 Charles Street, Cambridge, MA 02141, USA  
On Mar 16, 2000 this sequence version replaced gi:6446801.  
All repeats were identified using RepeatMasker:  
Smit, A.F.A. & Green, P. (1996-1997)  
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

----- Genome Center  
Center: Whitehead Institute/ MIT Center for Genome Research  
Center code: WIBR  
Web site: <http://www-seq.wi.mit.edu>  
Contact: [sequence\\_submissions@genome.wi.mit.edu](mailto:sequence_submissions@genome.wi.mit.edu)  
----- Project Information  
Center project name: L641  
Center clone name: 71\_A\_1  
----- Summary Statistics  
Sequencing vector: M13; M77815; 100% of reads  
Sequencing vector: Plasmid; n/a; 0.0 f% of reads  
0.273224043715847Chemistry: Dye-primer-amersham; 9% of reads  
Chemistry: Dye-terminator Big Dye; 91% of reads  
Assembly program: Phrap; version 0.960731  
Consensus quality: 147496 bases at least Q40  
Consensus quality: 154226 bases at least Q30  
Consensus quality: 156653 bases at least Q20  
Insert size: 137000; agarose-1p

gene expression with the option of cell growth control  
 Patent: WO 0065074-A 2 02-NOV-2000;  
 Gesellschaft für Biotechnologische Forschung mbH (GBF) (; DE)  
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 Db 61 AAATCAGATC 70

AC032033 69861 bp DNA linear HTG 03-APR-2000  
 Homo sapiens chromosome 18 clone RP11-392F20 map 18, LOW-PASS  
 SEQUENCE SAMPLING.

AC032033  
 AC032033.1 GI:7387453  
 HTG: HTGS.PHASE0.

SOURCE  
 Homo sapiens.

ORGANISM  
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 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE  
 1 (bases 1 to 69861)

AUTHORS  
 Birren,B., Linton,L., Nusbaum,C. and Lander,E.

TITLE  
 Homo sapiens chromosome 18, clone RP11-392F20

REFERENCE  
 2 (bases 1 to 69861)

AUTHORS  
 Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,  
 Anderson,S., Baldwin,J., Barna,N., Bastien,V., Beda,E.,  
 Boguslavskiy,L., Boukhgalter,B., Brown,A., Burkett,G.,  
 Campioniano,A., Castle,A., Choepel,Y., Colangelo,M., Collins,S.,  
 Collymore,A., Cooke,P., DeArelano,K., Dewar,K., Diaz,J.S.,  
 Dodge,S., Domino,M., Doyle,M., Ferreira,P., FitzHugh,W., Gage,D.,  
 Galagan,J., Gardyna,S., Glnde,S., Goyette,M., Graham,L.,  
 Grand-Pierre,N., Grant,G., Hagos,B., Heaford,A., Horton,L.,  
 Howland,J.C., Iliev,I., Johnson,R., Jones,C., Kann,L., Karatas,A.,  
 Klein,J., LaRocque,K., Lamazares,R., Landers,T., Lehoczy,J.,  
 Levine,R., Lieu,C., Liu,G., Locke,K., Macdonald,P., Margulis,N.,  
 McCarthy,M., McEwan,P., McGurk,A., McKernan,K., McPheeters,R.,  
 Melchior,J., Meneus,L., Mihova,T., Miranda,C., Mlenga,V., Morrow,J.,  
 Murphy,T., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,  
 O'Neill,D., Ollivar,T.M., Oliver,J., Peterson,K., Pierre,N.,  
 Pisani,C., Pollara,V., Raymond,C., Riley,R., Rogov,P., Rothman,D.,  
 Roy,A., Santos,R., Schauer,S., Severy,P., Spencer,B.,  
 Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,  
 Tesfaye,S., Theodore,J., Tirrell,A., Travers,M., Trigilio,J.,  
 Vassiliev,H., Viel,R., Vo,A., Willson,B., Wu,X., Wyman,D., Ye,W.J.,  
 Young,G., Zainoun,J., Zimmer,A. and Zody,M.

Direct Submission

Submitted (03-APR-2000) Whitehead Institute/MIT Center for Genome

Research, 320 Charles Street, Cambridge, MA 02141, USA

All repeats were identified using RepeatMasker:

Smit, A.F.A. & Green, P. (1996-1997)

http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: http://www.seq.wi.mit.edu

Contact: sequence\_submissions@genome.wi.mit.edu

----- Project Information  
 Center project name: L9119  
 Center clone name: 392\_F\_20

\* NOTE: This record contains 74 individual  
 \* sequencing reads that have not been assembled into  
 \* contigs. Runs of N are used to separate the reads  
 \* and the order in which they appear is completely  
 \* arbitrary. Low-pass sequence sampling is useful for  
 \* identifying clones that may be gene-rich and allows  
 \* overlap relationships among clones to be deduced.  
 \* However, it should not be assumed that this clone  
 \* will be sequenced to completion. In the event that  
 \* the record is updated, the accession number will  
 \* be preserved.

\* 1 848: contig of 848 bp in length  
 \* 849 948: gap of 100 bp  
 \* 949 1814: contig of 866 bp in length  
 \* 1815 1914: gap of 100 bp  
 \* 1915 2781: contig of 867 bp in length  
 \* 2782 2881: gap of 100 bp  
 \* 2882 3726: contig of 845 bp in length  
 \* 3727 3826: gap of 100 bp  
 \* 3827 4682: contig of 856 bp in length  
 \* 4683 4782: gap of 100 bp  
 \* 4783 5612: contig of 830 bp in length  
 \* 5613 5712: gap of 100 bp  
 \* 5713 6560: contig of 848 bp in length  
 \* 6561 6660: gap of 100 bp  
 \* 6661 7508: contig of 848 bp in length  
 \* 7509 7608: gap of 100 bp  
 \* 7609 8438: contig of 830 bp in length  
 \* 8439 8538: gap of 100 bp  
 \* 8539 9375: contig of 837 bp in length  
 \* 9376 9475: gap of 100 bp  
 \* 9476 10341: contig of 866 bp in length  
 \* 10342 10441: gap of 100 bp  
 \* 10442 11277: contig of 836 bp in length  
 \* 11278 11377: gap of 100 bp  
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 \* 12219 12318: gap of 100 bp  
 \* 12319 13169: contig of 851 bp in length  
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 \* 13270 13976: contig of 707 bp in length  
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 \* 14932 15031: gap of 100 bp  
 \* 15032 15872: contig of 841 bp in length  
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 \* 15973 16802: contig of 830 bp in length  
 \* 16803 16902: gap of 100 bp  
 \* 16903 17723: contig of 821 bp in length  
 \* 17724 17823: gap of 100 bp  
 \* 17824 18674: contig of 851 bp in length  
 \* 18675 18774: gap of 100 bp  
 \* 18775 19632: contig of 858 bp in length  
 \* 19633 19732: gap of 100 bp  
 \* 19733 20580: contig of 848 bp in length  
 \* 20581 20680: gap of 100 bp  
 \* 20681 21532: contig of 852 bp in length  
 \* 21533 21632: gap of 100 bp  
 \* 21633 22477: contig of 845 bp in length  
 \* 22478 22577: gap of 100 bp  
 \* 22578 23424: contig of 847 bp in length  
 \* 23425 23524: gap of 100 bp  
 \* 23525 24327: contig of 803 bp in length  
 \* 24328 24427: gap of 100 bp  
 \* 24428 25243: contig of 816 bp in length  
 \* 25244 25343: gap of 100 bp  
 \* 25344 26186: contig of 843 bp in length  
 \* 26187 26286: gap of 100 bp  
 \* 26287 27153: contig of 867 bp in length  
 \* 27154 27253: gap of 100 bp

GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: March 5, 2003, 22:29:16 ; Search time 385.676 Seconds  
(without alignments)  
5282.140 Million cell updates/sec

Title: us-09-980-548-2  
Perfect score: 70  
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Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues  
Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000  
Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

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- 2: gb\_htg.\*
- 3: gb\_in.\*
- 4: gb\_om.\*
- 5: gb\_ov.\*
- 6: gb\_pat.\*
- 7: gb\_ph.\*
- 8: gb\_pl.\*
- 9: gb\_pr.\*
- 10: gb\_ro.\*
- 11: gb\_sts.\*
- 12: gb\_sy.\*
- 13: gb\_un.\*
- 14: gb\_vl.\*
- 15: em\_ba.\*
- 16: em\_fun.\*
- 17: em\_hum.\*
- 18: em\_in.\*
- 19: em\_mu.\*
- 20: em\_om.\*
- 21: em\_or.\*
- 22: em\_ov.\*
- 23: em\_pat.\*
- 24: em\_ph.\*
- 25: em\_pl.\*
- 26: em\_ro.\*
- 27: em\_sts.\*
- 28: em\_un.\*
- 29: em\_vl.\*
- 30: em\_htg\_hum.\*
- 31: em\_htg\_inv.\*
- 32: em\_htg\_other.\*
- 33: em\_htg\_mus.\*
- 34: em\_htg\_pln.\*
- 35: em\_htg\_rod.\*
- 36: em\_htg\_mam.\*
- 37: em\_htg\_vrt.\*
- 38: em\_sy.\*
- 39: em\_htgo\_hum.\*
- 40: em\_htgo\_mus.\*
- 41: em\_htgo\_other.\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query % Match	Length	ID	Description
1	70	100.0	70	AX040910	Sequence
2	31.6	45.1	69861	2	AC032033 Homo sapi
3	31.6	45.1	161198	2	AC015867 Homo sapi
4	31.6	45.1	174396	9	AC104564 Homo sapi
5	30.6	43.7	13943	10	AF481734 Mus muscu
6	30.6	43.7	196360	2	AC098508 Rattus no
7	30.6	43.7	270795	2	AC015335 Mus muscu
8	30	42.9	62795	2	AC022602 Homo sapi
9	30	42.9	148555	2	AP004711 Homo sapi
10	30	42.9	160883	9	AC021498 Homo sapi
11	30	42.9	197743	10	AL591711 Mouse DNA
12	29.6	42.3	90747	2	AC011975 Homo sapi
13	29.2	41.7	110121	9	AL590989 Human DNA
14	29.2	41.7	176775	2	AC016978 Homo sapi
15	29	41.4	165373	9	AC067957 Homo sapi
16	29	41.4	173874	2	AC067791 Homo sapi
17	28.8	41.1	61802	2	OSIG00028
18	28.8	41.1	141128	8	OSJN00139
19	28.6	40.9	66317	2	AC110321
20	28.6	40.9	213937	10	AL672128 Mouse DNA
21	28.4	40.6	264461	2	AC130804
22	28.2	40.3	70522	9	HS130491
23	28.2	40.3	78603	9	HS1188A21
24	28.2	40.3	176774	2	AP002519
25	28	40.0	31976	9	AL596107
26	28	40.0	120724	9	CNS05TCR
27	28	40.0	144223	9	AC062015
28	28	40.0	145458	2	AC069591
29	28	40.0	197659	9	AC009245
30	27.8	39.7	125590	9	AC002382
31	27.8	39.7	141905	2	AC110101
32	27.8	39.7	161831	2	AC073078
33	27.8	39.7	163567	2	AC120955
34	27.8	39.7	180318	2	AC109991
35	27.8	39.7	186862	2	AC110483
36	27.8	39.7	214817	2	AC127366
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40	27.6	39.4	136641	2	AC114327
41	27.6	39.4	168703	4	AC091759
42	27.6	39.4	172884	2	AC113567
43	27.6	39.4	184234	2	AC091758
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ALIGNMENTS

RESULT 1	AX040910	Sequence 2 from Patent WO0065074.	70 bp	DNA	linear	PAT 23-NOV-2000
LOCUS	AX040910	Sequence 2 from Patent WO0065074.				
DEFINITION	AX040910					
ACCESSION	AX040910					
VERSION	AX040910.1	GI:11340532				
KEYWORDS						
SOURCE	Mesocricetus sp.					
ORGANISM	Mesocricetus sp.					
	Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;					
	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Cricetinae;					
	Mesocricetus.					
REFERENCE	1. (bases 1 to 70)					
AUTHORS	Mueller, P., Geserick, C., Schroeder, K. and Hauser, H.					
TITLE	Promoter-transactivator system for inducible high-level mammalian					

Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y. Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. *Genome Res.* 10 (10), 1617-1630 (2000)

wagi, K., Fujiwaka, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.

RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. *Genome Res.* 10 (11), 1757-1771 (2000)

Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara, Y. and Hayashizaki, Y.

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. *Genome Res.* 11 (2), 281-289 (2001)

Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.

e mouse tissues.

e mouse tissues.  
Location/Qualifiers  
1. .535

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ERG 174

/note="Vector: pSPORT1; Site.1: SalI; Site.2: NotI; This clone is among a rearranged set of 15,247 clones from 11 embryo cDNA libraries (including preimplantation stage embryos from unfertilized egg to blastocyst, embryonic part of E7.5 embryos, extraembryonic part of E7.5 embryos, and E12.5 female mesonephros/gonad) and one newborn ovary cDNA library. Average insert size 1.5 kb. All source libraries are cloned unidirectionally with Oligo(dT)-Not primers. References include: (1) Genome-wide expression profiling of mid-gestation placenta and embryo using a 15,000 mouse developmental cDNA microarray, 2000, Proc. Natl. Acad. Sci. U S A, 97: 9127-9132; (2) Large-scale cDNA analysis reveals phased gene expression patterns during preimplantation mouse development, 2000, Development, 127: 1737-1749; (3) Genome-wide mapping of unselected transcripts from extraembryonic tissue of 7.5-day mouse embryos reveals enrichment in the t-complex and under-representation on the X chromosome, 1998, Hum Mol Genet 7: 1967-1978."

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Db 142 CAGGATATCTGTGTCAAGCACCTGGCCCGCTCAGGCGCCAGAACAGATGGTAC 198

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LOCUS      BB756171
DEFINITION BB756171 RIKEN full-length enriched, melanocyte Mus musculus cDNA
ACCESSION BB756171.1 GI:16202641
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SOURCE     house mouse.
ORGANISM   Mus musculus
REFERENCE   Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS     Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
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            Akimura,T., Arakawa,T., Carninci,P., Furuno,M., Hanagaki,T.,
            Hayatsu,N., Hiramoto,K., Hiraoka,T., Hirozane,T., Imotani,K., Ishii
            ,Y., Ito,M., Kawai,J., Kojima,Y., Konno,H., Kouda,M., Matsuyama,T.,
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            Shibata,K., Shinagawa,A., Shiraki,T., Sogabe,Y., Suzuki,H., Tagawa
            ,A., Takahashi,F., Takaku-Akahira,S., Tanaka,T., Tomaru,A., Toya,T.,
            Watahiki,A., Yasunishi,A., Muramatsu,M. and Hayashizaki,Y.
            RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura,T., et al.
            2001)
JOURNAL     Unpublished (2001)
COMMENT     Contact: Yoshihide Hayashizaki
            Laboratory for Genome Exploration Research Group, RIKEN Genomic
            Sciences Center(GSC), Yokohama Institute
            The Institute of Physical and Chemical Research (RIKEN)
            1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
            Tel: 81-45-503-9226
            Fax: 81-45-503-9216
            Email: genome-res@gsc.riken.go.jp
            URL: http://genome.gsc.riken.go.jp/
            Carninci,P., Shibata,Y., Hayatsu,N., Sugahara,Y., Shibata,K., Itoh

```

M., Konno,H., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.  
 Normalization and subtraction of cap-trapper-selected cDNAs to  
 prepare full-length cDNA libraries for rapid discovery of new  
 genes. Genome Res. 10 (10), 1617-1630 (2000)  
 wagi,K., Fujiwaki,S., Inoue,K., Togawa,Y., Izawa,M., Ohara,E.,  
 Watahiki,M., Yoneda,Y., Ishikawa,T., Ozawa,K., Tanaka,T., Matsura  
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 RIKEN integrated sequence analysis (RISA) system--384-format  
 sequencing pipeline with 384 multicapillary sequencer. Genome Res.  
 10 (11), 1757-1771 (2000)  
 Konno,H., Fukunishi,Y., Shibata,K., Itoh,M., Carninci,P., Sugahara  
 ,Y. and Hayashizaki,Y.  
 Computer-based methods for the mouse full-length cDNA  
 encyclopedia: real-time sequence clustering for construction of a  
 nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)  
 Please visit our web site (<http://genome.gsc.riken.go.jp>) for  
 further details.  
 e mouse tissues.

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ORIGIN
Query Match      21.9%; Score 59; DB 10; Length 506;
Best Local Similarity 67.2%; Pred. No. 2e-05;
Matches 119; Conservative 0; Mismatches 45; Indels 13; Gaps 2;

QY 1 GCTAGCTTAAGTACGCCATTTTCAGGCGATGGGAAATACATACTGAGATAGAGA 60
Db 72 GCTAGCTCAGTACGCCATTTTCAGGCGATGGGAAATACATACTGAGATAGAGA 131
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QY 109 CAGGATATCTGTGT-AAGCAGTTCTCTGCCCGCTCAGGCGCCAGAACAGATTGGAAC 164
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RESULT 14
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DEFINITION BB753609 NIH_MGC_114 Homo sapiens cDNA clone IMAGE:5198973 5',
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ACCESSION   BB753609
VERSION     BB753609.1 GI:15745187
KEYWORDS    EST.
SOURCE      human.
ORGANISM    Homo sapiens
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REFERENCE   1 (bases 1 to 787)
AUTHORS     NIH-MGC http://mgc.nci.nih.gov/.
TITLE       National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL     Unpublished (1999)
COMMENT     Contact: Robert Strausberg, Ph.D.
            Email: cagabp-r@mail.nih.gov
            Tissue Procurement: Life Technologies, Inc.

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**TITRE**

**Qy** 109 CAGGATATCTGGT - AAGCAGTTCTGCCCGCCGCTCAGGGCCAGAACAGTTGGAAC 164  
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**Db** 142 CAGGATATCTGGTCAAGCACCTGGGCCCGGCCTCAGGGCCAGAACAGATGGTAC 198

134 a 127 c 98 g 97 t

21.9%; Score 59; DB 10; Length 456;  
Similarity 67.2%; Pred. No. 1.9e-05;

Length 21.9%; Score 59; DB 10; Length 456;  
 Similarity 67.2%; Pred. No. 1.9e-05;  
 Conservative 0; Mismatches 45; Indels 13; Gaps 2;  
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 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||  
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Y. and Hayashizaki, Y. Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. *Genome Res.* 11 (2), 281-289 (2001). Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.

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Location/Qualifiers
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Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in
RIKEN, Division of Experimental Animal Research in Riken
contributed to prepare mouse tissues. 1st strand cDNA was
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GAGAGAGAGAGATCTCGAGTCTAATTAATTAATCTCCCCCCCCCCC 21'. cDNA was prepared by using trehalase thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. Second strand cDNA was prepared with the primer adapter of sequence [5',

134 a 126 c 98 g 97 t

h 21.9%; Score 59; DB 10; Length 455;  
 Similarity 67.2%; Pred. No. 1.9e-05;  
 19; Conservative 0; Mismatches 45; Indels 13; Gaps 2;  
  
 TAGCTTAAAGTAACGCCATTTTCGAAGCATGGGAAAAAATACATAACTCAGAAATACAGA 60  
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 TTAGCTCAGTAAGCGCCATTGTGCAAGGCATGA AAAAGTACCAGAGCTAGGTTCTCAA 80  
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||  
 TTTCAGATCAAGGTCAGGAACAGAAACAGGAGAA-----ATCGGCCAAA 108  
 || | || | || | || | || | || | || | || | || | || | || | || | || | || |  
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BB726927 BB726927  
BB726927.1 GI:16110202  
EST.  
house mouse.  
Mus musculus  
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Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
1 (bases 1 to 456)



Fax: 81-45-503-9216  
 Email: genome-res@gsc.riken.go.jp,  
 URL: http://genome.gsc.riken.go.jp/  
 Carninci, P., Shibata, Y., Muramatsu, M., Sugahara, Y., Shibata, K., Itoh  
 M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.  
 Normalization and subtraction of cap-trapper-selected cDNAs to  
 prepare full-length cDNA libraries for rapid discovery of new  
 genes. Genome Res. 10 (10), 1617-1630 (2000)  
 wagi, K., Fujiwaka, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E.,  
 Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura  
 S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and  
 Hayashizaki, Y.  
 RIKEN integrated sequence analysis (RISA) system--384-format  
 sequencing pipeline with 384 multicapillary sequencer. Genome Res.  
 10 (11), 1757-1771 (2000)  
 Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara  
 Y., and Hayashizaki, Y.  
 Computer-based methods for the mouse full-length cDNA  
 encyclopedia: real-time sequence clustering for construction of a  
 nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)  
 Please visit our web site (http://genome.gsc.riken.go.jp) for  
 further details.  
 e mouse tissues.

UNIGEN  
 : COUNT 130 a 120 c 93 g 96 t  
 Query Match 21.9%; Score 59; DB 10; Length 439;  
 Best Local Similarity 67.2%; Pred. No. 1.9e-05;  
 Matches 119; Conservative 0; Mismatches 45; Indels 13; Gaps 2;  
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 Db 66 AGTTTCAAGGAAGTTAGTTAAATAAATAAAGGCTGAATAATCTAGACAGAGGCCCAA 125  
 QY 109 CAGGATATCTGTGGT-AAGCACTTCTGCCCGCTCAGGCGCAAGCAAGCAAGTGTGGAC 164  
 Db 126 CAGGATATCTGTGGTCAAGCACTTGGGCGGCTCAGGCGCAAGCAAGCAAGTGTGGTAC 182

RESULT 9

BB670654  
 LOCUS  
 DEFINITION  
 ACCESSION  
 VERSION  
 KEYWORDS  
 SOURCE  
 ORGANISM  
 REFERENCE  
 AUTHORS  
 TITLE  
 JOURNAL  
 COMMENT

BB670654 445 bp mRNA linear EST 05-OCT-2001  
 BB670654 RIKEN full-length enriched, 17 days embryo head Mus  
 musculus cDNA clone 3322404N14 3', mRNA sequence.  
 BB670654  
 EST  
 BB670654.1 GI:15969875  
 house mouse.  
 Mus musculus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 Akimura, T., Hirakawa, T., Carninci, P., Furuno, M., Hanagaki, T.,  
 Hayatsu, N., Hiramoto, K., Hiraoka, T., Hirozane, T., Imotani, K., Ishii  
 Y., Ito, M., Kawai, J., Kojima, Y., Konno, H., Kouda, M., Matsuyama, T.,  
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 Watahiki, A., Yasunishi, A., Muramatsu, M. and Hayashizaki, Y., et al.  
 RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura, T., et al.  
 2001)

Unpublished (2001)  
 Contact: Yoshihide Hayashizaki  
 Laboratory for Genome Exploration Research Group, RIKEN Genomic  
 Sciences Center (GSC), Yokohama Institute  
 The Institute of Physical and Chemical Research (RIKEN)  
 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan  
 Tel: 81-45-503-9222  
 Fax: 81-45-503-9216  
 Email: genome-res@gsc.riken.go.jp,  
 URL: http://genome.gsc.riken.go.jp/  
 Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh  
 M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.  
 Normalization and subtraction of cap-trapper-selected cDNAs to  
 prepare full-length cDNA libraries for rapid discovery of new  
 genes. Genome Res. 10 (10), 1617-1630 (2000)  
 wagi, K., Fujiwaka, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E.,  
 Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura  
 S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and  
 Hayashizaki, Y.  
 RIKEN integrated sequence analysis (RISA) system--384-format  
 sequencing pipeline with 384 multicapillary sequencer. Genome Res.  
 10 (11), 1757-1771 (2000)  
 Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara  
 Y., and Hayashizaki, Y.  
 Computer-based methods for the mouse full-length cDNA  
 encyclopedia: real-time sequence clustering for construction of a  
 nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)  
 Please visit our web site (http://genome.gsc.riken.go.jp) for  
 further details.  
 e mouse tissues.

FEATURES  
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Location/Qualifiers  
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 /strain="C57BL/6J"  
 /db\_xref="taxon:10090"  
 /clone="3322404N14"  
 /clone\_lib="RIKEN full-length enriched, 17 days embryo  
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 /sex="mixed"  
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/note="Site\_1: Sali; Site\_2: BamHI; cDNA library was  
 prepared and sequenced in Mouse Genome Encyclopedia  
 Project of Genome Exploration Research Group in Riken  
 Genomic Sciences Center and Genome Science Laboratory in  
 RIKEN. Division of Experimental Animal Research in Riken  
 contributed to prepare mouse tissues. 1st strand cDNA was  
 primed with a primer [5'  
 GAGAGAGAGAGATCCAGAGCTCTTTTTTTTTTTTNN 3'], cDNA was  
 prepared by using trehalose thermo-activated reverse

Brazil

Tel: +55-11-2704922  
Fax: +55-11-2707001  
Email: asimpson@ludwig.org.br

This sequence was derived from the FAPESP/LICR Human Cancer Genome project. This entry can be seen in the following URL  
(http://www.ludwig.org.br/scripts/gethtml2.pl?tl1=RC3&tl2=RC5-BT0604-260100-011-C11&t3=2000-01-26&t4=1)

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musculus cDNA clone E970034P16 3', mRNA sequence.  
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 EST.  
 house mouse.  
 Mus musculus  
 Eukaryota; Metazoa: Chordata: Vertebrata; Euteleostomi;  
 Mammalia; Eutheria: Rodentia: Sciurognathi; Muridae; Murinae; Mus.  
 1 (bases 1 to 439)  
 Akimura,T., Arakawa,T., Carninci,P., Furuno,M., Hanagaki,T.,  
 Hayatsu,N., Hiramoto,K., Hiraoka,T., Hirozane,T., Imotani,K., Ishii,  
 Y., Ito,M., Kawai,J., Kojima,Y., Konno,H., Kouda,M., Matsuyama,T.,  
 Nakamura,M., Nishi,K., Nomura,K., Numasaki,R., Okazaki,Y., Okido,T.,  
 Saito,R., Sakai,C., Sakai,K., Sakazume,N., Sasaki,D., Sato,K.,  
 Shibata,K., Shinagawa,A., Shiraki,T., Sogabe,Y., Suzuki,H., Tagawa,  
 A., Takahashi,F., Takaku-Akahira,S., Tanaka,T., Tomaru,A., Taya,T.,  
 Watahiki,A., Yasunishi,A., Muramatsu,M. and Hayashizaki,Y.  
 RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura,T., et al.  
 2001)  
 Unpublished (2001).  
 Contact: Yoshihide Hayashizaki  
 Laboratory for Genome Exploration Research Group, RIKEN Genomic  
 Sciences Center(GSC), Yokohama Institute  
 The Institute of Physical and Chemical Research (RIKEN)  
 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan  
 Tel: 81-45-503-9222

Query Match	22.88;	Score 61.6;	DB 10;	Length 444;
Best Local Similarity	68.24;	Pred. No. 3.9e-06;		
Matches 120;	Conservative	0;	Mismatches 44;	Indels 12;
Gaps	2;			
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QY	110	AGGATATCTGTGTT-AGCAGTTCCTCGCCGCTCAGGCGCCAGCAAGACAGTTGGGAAC 164		
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ACCESSION	BB835008			
VERSION	BB835008.1	GI:17013251		
KEYWORDS	EST.			
SOURCE	house mouse.			
ORGANISM	Mus musculus			
REFERENCE	1 (bases 1 to 446)			
AUTHORS	Akimura, T., Arakawa, T., Carninci, P., Furuno, M., Hanagaki, T., Hayatsu, N., Hiramoto, K., Hiraoka, T., Hirozane, T., Imotani, K., Ishii, Y., Ito, M., Kawai, J., Kojima, Y., Konno, H., Kouda, M., Matsuyama, T., Nakamura, M., Nishi, K., Nomura, K., Numasaki, R., Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K., Sakazume, N., Sasaki, D., Sato, K., Shibata, K., Shinagawa, A., Shiraki, T., Sogabe, Y., Suzuki, H., Tagawa, A., Takahashi, F., Takaku-Akahira, S., Tanaka, T., Tomaru, A., Toya, T., Watahiki, A., Yasunishi, A., Muramatsu, M. and Hayashizaki, Y.			
TITLE	RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura, T., et al. 2001)			
JOURNAL	Unpublished (2001)			
COMMENT	Contact: Yoshihide Hayashizaki Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute The Institute of Physical and Chemical Research (RIKEN) 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222 Fax: 81-45-503-9216 Email: genome-res@gsc.riken.go.jp/ URL: http://genome.gsc.riken.go.jp/ Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y. Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000) wagi, K., Fujiwaka, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y. RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000) Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara Y. and Hayashizaki, Y. Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001) Please visit our web site (http://genome.gsc.riken.go.jp) for further details. e mouse tissues.			
FEATURES	Location/Qualifiers			
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	/strain="C57BL/6J"			
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ACCESSION    BF101077
VERSION      BF101077.1  GI:10883603
KEYWORDS     EST.
SOURCE       house mouse.
ORGANISM     Mus musculus.

REFERENCE    1 (bases 1 to 1240)
AUTHORS      NIH-MGC http://mgc.nci.nih.gov/
TITLE        National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL      Unpublished (1999)
COMMENT      Contact: Robert Strausberg, Ph.D.
            Email: cgapbs-re@mail.nih.gov
            Tissue Procurement: Gilbert Smith, Ph.D.
            cDNA Library Preparation: Life Technologies, Inc.
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: Incyte Genomics, Inc.
            Clone distribution: MGC clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            http://image.llnl.gov
            Plate: LLM9184 row: b column: 12
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                Library constructed by Life Technologies. Investigator
                providing samples: Gilbert Smith, NIH"
                providing samples: Gilbert Smith, NIH"
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Matches 120; Conservative 0; Mismatches 44; Indels 11; Gaps 2;

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ACCESSION    AZ804164
VERSION      AZ804164.1  GI:12956487
KEYWORDS     GSS.
SOURCE       house mouse.
ORGANISM     Mus musculus.

REFERENCE    1 (bases 1 to 666)
AUTHORS      Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,

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Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
and Wright,D.,Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
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                was hydrodynamically sheared by repeated passage through a
                0.005 inch orifice at constant velocity. The sheared DNA
                was blunt end-repaired with T4 DNA polymerase and T4
                polynucleotide kinase. Adaptor oligonucleotides were
                ligated to the blunt ends in high molar excess. The
                adapted DNA was purified and size-selected for a 9.5 to
                10.5 kb range using preparative agarose gel
                electrophoresis. Vector DNA was prepared from a derivative
                of pWD42 (gil4732114|gb|AF129072.1), a copy-number
                inducible derivative of plasmid R1. The vector was ligated
                with adaptors complementary to the insert adaptors and
                purified. The sheared, adapted mouse DNA was annealed to
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                chemically-competent E. coli XL10-Gold (Stratagene) cells
                and selected for ampicillin resistance."
                and selected for ampicillin resistance."
BASE COUNT   141 a 166 c 148 g 211 t
ORIGIN
Query Match      22.9%; Score 61.8; DB 17; Length 666;
Best Local Similarity 58.1%; Pred. No. 3.7e-06;
Matches 155; Conservative 0; Mismatches 97; Indels 15; Gaps 2;

Qy 1  GCTAGCTTAAGTAACGCCATTTTGCAGGCATGGGAAATACATACACTGAGATAGAGA 60
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Qy 121  GGTAAAGCAGTTCTCCCGCTCAGGCGCCCAAGAACAGTTGGAACAGGAGAAATGGGCCAA 180
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Qy 181  ACAGGATATCTGTGGTAAGCAGATTCTCTCCCGCTCAGGCGCCCAAGAACAGATGGTCCCA 240
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```

Tel: 81-45-503-9222  
Fax: 81-45-503-9216

Email: genome-res@gsr.riken.go.jp,  
URL: http://genome.gsc.riken.go.jp/  
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh  
M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.  
Normalization and subtraction of cap-trapper-selected cDNAs to  
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S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and  
Hayashizaki, Y.

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Y. and Hayashizaki, Y.

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Please visit our web site (http://genome.gsc.riken.go.jp) for  
further details.

e mouse tissues.

.FURES  
Source

Location/Qualifiers  
1. 538

/organism="Mus musculus"

/strain="C57BL/6J"

/db\_xref="taxon:10090"

/clone="G270003K01"

/cell\_type="melanocyte"

/note="pooled tissues; (tissue\_type=cerebellum,  
dev\_stage=16 days neonate, sex=mixed)

(tissue\_type=cerebellum, dev\_stage=0 day neonate,  
sex=mixed), (tissue\_type=hippocampus, dev\_stage=adult,  
sex=adult), (tissue\_type=whole body, dev\_stage=9 days  
embryo, sex=mixed), (tissue\_type=liver, dev\_stage=13 days  
embryo, sex=mixed)"

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Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.  
1 (bases 1 to 616)

REFERENCE

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Unpublished (2000)

Contact: Robert B. Weiss

University of Utah Genome Center

University of Utah

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT

84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: dduenne@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Plate: 0266 row: C column: 09

Seq primer: CGTTGTAAACGACGCCAGT

Class: plasmid ends

High quality sequence stop: 616.

Location/Qualifiers

1. 616

/organism="Mus musculus"

/strain="C57BL/6J"

/db\_xref="taxon:10090"

/clone="UUGC2M0266C09"

/clone\_lib="Mouse 10kb plasmid UUGC2M library"

/sex="Female"

/lab\_host="E. coli strain XL10-Gold, T1-resistant, F."

/note="Vector: PWD42nv: Purified genomic DNA from M.  
musculus C57BL/6J (female) was obtained from the Jackson  
Laboratory Mouse DNA Resource  
(http://www.jax.org/resources/documents/dnares/). The DNA  
was hydrodynamically sheared by repeated passage through a  
0.005 inch orifice at constant velocity. The sheared DNA  
was blunt end-repaired with T4 DNA polymerase and T4  
polynucleotide kinase. Adaptor oligonucleotides were  
ligated to the blunt ends in high molar excess. The  
adapted DNA was purified and size-selected for a 9.5 to  
10.5 kb range using preparative agarose gel  
electrophoresis. Vector DNA was prepared from a derivative  
of pWD42 (gil4732114|gb|AF129072.1), a copy-number  
inducible derivative of plasmid R1. The vector was ligated  
with adaptors complementary to the insert adaptors and  
purified. The sheared, adapted mouse DNA was annealed to  
adapted vector DNA, and transformed into  
chemically-competent E. coli XL10-Gold (Stratagene) cells  
and selected for ampicillin resistance."

BASE COUNT

180 a 151 c 141 g 144 t

ORIGIN

Query Match

Best Local Similarity

Matches 120; Conservative

0; Mismatches 44; Indels 11; Gaps 2;

QY 1 GCTAGCTTAAGTAACCCATTTTTCGAAGCATGGGAAATAACATCACTGAGAATAGAGA 60

Db 116 GCTAACTGCAGTACGCCATCTTTCGAAGCATGGGAAATAACATCACTGAGAATAGAGA 175

QY 61 AGTTCAGATCAAGGTCAGGACAGAAACAGGAGAAATAT-----GGGCAACACAGG 112

176 AAAACACAGACAGGAGTACAGAGAGCTGGAAAGTACCGGAGTAGGGCCCAACACAGG 235

QY 113 ATATCTGTGGTAAGCAGTCTCTCCCGCTCAGGGCCCAAGACAGTTGCAACAGGAGAAAT 172

Db 236 ATATCTGTGGTAAGCAGTCTCTCCCGCTCAGGGCCCAAGACAGTTGCAACAGGAGAAAT 295

QY 173 TG 174

Db 296 AG 297

RESULT 2

AZ984553

LOCUS

DEFINITION

2M0266C09F Mouse 10kb plasmid UUGC2M library Mus musculus genomic

clone UUGC2M0266C09 F, DNA sequence.

ACCESSION

AZ984553

VERSION

AZ984553.1

GI:13855780

KEYWORDS

GSS.

SOURCE

house mouse.

ORGANISM

Mus musculus

616 bp DNA linear

GSS 27-APR-2001

GenCore version 5.1.3  
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model  
Run on: March 5, 2003, 22:43:11 ; Search time 2036.09 Seconds  
(without alignments)  
2147.640 Million cell updates/sec

Title: US-09-980-548-1  
Perfect score: 270  
Sequence: 1 gctagtttaagtaagccat.....cgccctcagcagttttctaga 270

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

-eached: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : EST:\*

- 1: em\_estba:\*
- 2: em\_esthum:\*
- 3: em\_estin:\*
- 4: em\_estmu:\*
- 5: em\_estov:\*
- 6: em\_estpl:\*
- 7: em\_estro:\*
- 8: em\_htc:\*
- 9: gb\_estl:\*
- 10: gb\_est2:\*
- 11: gb\_hic:\*
- 12: gb\_est3:\*
- 13: gb\_est4:\*
- 14: gb\_est5:\*
- 15: em\_estfun:\*
- 16: em\_estom:\*
- 17: gb\_gss:\*
- 18: em\_gss\_hum:\*
- 19: em\_gss\_inv:\*
- 20: em\_gss\_pln:\*
- 21: em\_gss\_vrt:\*
- 22: em\_gss\_fun:\*
- 23: em\_gss\_mam:\*
- 24: em\_gss\_mus:\*
- 25: em\_gss\_other:\*
- 26: em\_gss\_pro:\*
- 27: em\_gss\_rod:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	63.2	23.4	538	10	BB751692
2	62.6	23.2	616	17	AZ984553
3	62.6	23.2	1240	12	BF101077
4	61.8	22.9	666	17	AZ804164
5	61.6	22.8	444	10	BB750356
6	61.6	22.8	446	10	BB835008

c	7	59.4	22.0	284	10	AW750456
	8	59	21.9	439	10	BB733109
	9	59	21.9	445	10	BB670654
	10	59	21.9	455	10	BB670350
	11	59	21.9	456	10	BB726927
	12	59	21.9	456	10	BB733115
	13	59	21.9	506	10	BB756171
	14	58.8	21.8	787	13	BI753609
	15	58.6	21.7	535	10	BB762345
	16	58.6	21.7	537	10	BB762595
c	17	58.6	21.7	584	13	BI794183
	18	58.6	21.7	596	13	BI677171
	19	58.6	21.7	644	10	BB795130
	20	58.6	21.7	649	10	BB795436
	21	58.6	21.7	650	10	BB794939
	22	58.6	21.7	660	10	BB794906
	23	58.6	21.7	662	10	BB795586
	24	58.6	21.7	663	10	BB794704
	25	58.6	21.7	666	10	BB795529
	26	58.6	21.7	680	10	BB794600
	27	58.6	21.7	680	10	BB794643
	28	58.6	21.7	687	10	BB794760
	29	58.6	21.7	688	10	BB794665
	30	58.6	21.7	691	10	BB794587
	31	57.6	21.3	209	9	AA666591
	32	57.6	21.3	209	9	AA146207
	33	57.6	21.3	210	9	AA655589
	34	57.6	21.3	210	9	AA655895
	35	57.6	21.3	210	9	AA684134
	36	57.6	21.3	210	9	AA122917
	37	57.6	21.3	269	9	AA139125
	38	57.6	21.3	444	10	BB831589
	39	57.6	21.3	426	12	BF162178
	40	57	21.1	726	10	BB786738
	41	57	21.1	435	10	BB821235
	42	57	21.1	444	10	BB788783
	43	57	21.1	449	10	BB815029
	44	57	21.1	461	10	BB833539
	45	57	21.1	525	10	BB762415

ALIGNMENTS

RESULT 1  
BB751692  
LOCUS BB751692 RIKEN full-length enriched, melanocyte Mus musculus CDNA  
DEFINITION clone G270003K01 3', mRNA sequence.  
ACCESSION BB751692  
VERSION BB751692  
KEYWORDS EST.  
SOURCE house mouse.  
ORGANISM Mus musculus  
REFERENCE 1 (bases 1 to 538)  
AUTHORS Akimura,T., Hiramoto,K., Hiraoka,T., Hirozane,T., Imotani,K., Ishii Hayatsu,M., Kawai,J., Kojima,Y., Konno,H., Kouda,M., Matsuyama,T., Nakamura,M., Nishi,K., Nomura,K., Numasaki,R., Okazaki,Y., Okido,T., Saito,R., Sakai,C., Sakai,K., Sakazume,N., Sasaki,D., Sato,K., Shibata,K., Shinagawa,A., Shiraki,T., Sogabe,Y., Suzuki,H., Tagawa,A., Takahashi,F., Takaku-Akahira,S., Tanaka,T., Tomaru,A., Toya,T., Watahiki,A., Yasunishi,A., Muramatsu,M. and Hayashizaki,Y.  
RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura,T., et al. 2001)

JOURNAL Unpublished (2001)  
COMMENT Contact: Yoshihide Hayashizaki  
Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center(GSC), Yokohama Institute  
The Institute of Physical and Chemical Research (RIKEN)  
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan

CC proteins in mammalian tissue culture host cells, including rat fibroblast  
CC cells, bovine kidney cells and human kidney cells. The present sequence  
CC is cytomegalovirus (CMV) MN14 vector used in the invention. The vector  
CC comprises the following elements: CMV promoter, MN14 antibody heavy  
CC chain signal peptide, MN14 antibody heavy chain, encephalomyocarditis  
CC virus (ECMV) IRES -bovine alpha-lactalbumin signal peptide, MN14 antibody  
CC light chain and 3' moloney murine leukemia virus (MoMuLV) LTR.

XX Sequence 4207 BP; 1030 A; 1202 C; 1058 G; 917 T; 0 other;

Query Match 75.1%; Score 202.8; DB 24; Length 4207;  
Best Local Similarity 95.6%; Pred. No. 1.1e-55;  
Matches 262; Conservative 0; Mismatches 7; Indels 5; Gaps 5;

Oy 1 GCTAGCTTAAGTAAGCCCATTTTGGCAAGGCATGGGAAATAATACATACTAGAGATAGAGA 60

Db 3644 GCTAGCTTAAGTAAGCCCATTTTGGCAAGGCAT-GGAAAAATACATACTAGAGATAGAGA 3702

Oy 61 AGTTCAGATCAAGGTCAGGAACAGA-GAAACAGGAGAAATATGGCCCAACAGGATATCTG 119

Db 3703 AGTTCAGATCAAGGTCAGGAACAGATGGAAACAGCTGAATATGGCCCAACAGGATATCTG 3762

Oy 120 TGGTAAGCAGTTCTGCCCC-GCTCAGGGCCCAAGAACAGTTGGAAACAGGAGAAAT-TGGGC 177

Db 3763 TGGTAAGCAGTTCTGCCCCGCTCAGGGCCCAAGAACAGATGGAACAGCTGAATATGGGC 3822

Oy 178 CAACAGGATATCTGTGGTAAGCAGTTCTGCCCC-GCTCAGGGCCCAAGAACAGATGGTC 236

Db 3823 CAACAGGATATCTGTGGTAAGCAGTTCTGCCCCGCTCAGGGCCCAAGAACAGATGGTC 3882

Oy 237 CCCAGATCGGTCGCCGCCCTCAGCAGTTCTAGA 270

Db 3883 CCCAGATCGGTCGCCGCCCTCAGCAGTTCTAGA 3916

Search completed: March 5, 2003, 22:49:51  
Job time : 277.478 secs

two integrated integrating vectors. The integrating vectors comprise at least one exogenous gene operably linked to a promoter. The host cell is useful for producing a desired protein and for comparing protein functions. The host cell comprises a reporter gene which is from gene fluorescent protein, luciferase, beta-galactosidase and beta-lactamase, and the assaying step further comprises detecting a signal from the reporter gene. The desired protein includes proteins for pharmaceutical, industrial, diagnostic and other purposes. The host cells is useful for indirectly detecting the expression of a desired protein, comprising providing the host cell transfected with a vector encoding a polycistronic sequence comprising a signal protein and a desired protein operably linked by an internal ribosome entry site (IRES), and culturing the host cell under suitable conditions so that the signal protein and the desired protein is produced, where the presence of signal protein indicates the presence of desired protein. The present sequence is cytomagalovirus (CMV) MN14 vector used in the invention. The vector comprises the following elements: CMV promoter, MN14 antibody heavy chain signal peptide, MN14 antibody heavy chain, encephalomyocarditis virus (ECMV) IRES -bovine alpha-lactalbumin signal peptide, MN14 antibody light chain and 3' moloney murine leukemia virus (MoMuLV) LTR.

Sequence 4207 BP; 1030 A; 1202 C; 1058 G; 917 T; 0 other;

very Match 75.1%; Score 202.8; DB 24; Length 4207;  
 JEST Local Similarity 95.6%; Pred. No. 1.1e-55;  
 Matches 262; Conservative 0; Mismatches 7; Indels 5; Gaps 5;

Qy 1 GCTAGCTTAAGTAACGCCATTTTGGCAAGCATGGGAAATAATACATAACTGAGAATAGAGA 60  
 |||||||  
 Db 3644 GCTAGCTTAAGTAACGCCATTTTGGCAAGCAT-GGAAAAAATACATAACTGAGAATAGAGA 3702  
 |||||||  
 Qy 61 AGTTCAGATCAAGGTCAGGAACAGA-GAACAGAGAGATATGGCCAAACAGGATATCTG 119  
 |||||||  
 Db 3703 AGTTCAGATCAAGGTCAGGAACAGATGGAACAGCTGAATATGGCCAAACAGGATATCTG 3762  
 |||||||  
 Qy 120 TGGTAAGCAGTTCTTCGCCCC-GCTCAGGCGCCAAAGAACAGTGGCAAGAGAGAT-TGGGCG 177  
 |||||||  
 Db 3763 TGGTAAGCAGTTCTTCGCCCCGCTCAGGCGCCAAAGAACAGATGGAACAGCTGAATATGGGCG 3822  
 |||||||  
 Qy 178 CAACAGCATATCTGTGTAACAGTTCCTGCCCC-GCTCAGGCGCCAAAGAACAGATGGTC 236  
 |||||||  
 Db 3823 CAACAGCATATCTGTGTAACAGTTCCTGCCCCGCTCAGGCGCCAAAGAACAGATGGTC 3882  
 |||||||  
 Qy 237 CCCAGATCGGTCGCCGCCCTCAGCAGTTTCTTAGA 270  
 |||||||  
 Db 3883 CCCAGATCGGTCGCCGCCCTCAGCAGTTTCTTAGA 3916  
 |||||||

## RESULT 15

AAD28306  
 ID AAD28306 standard; DNA; 4207 BP.

AAD28306;

DT 22-APR-2002 (first entry)

DE Cytomegalovirus (CMV) MN14 vector.

KW Regulatory element; vector; erythropoietin; growth hormone; insulin;  
 KW immunoglobulin; bone morphogenetic protein; interferon; interleukin;  
 KW superoxide dismutase; T-cell receptor; surface membrane protein; CMV;  
 KW viral antigen; transport protein; addressin; regulatory protein; ECMV;  
 KW encephalomyocarditis virus; internal ribosome entry site; IRES; bovine;  
 KW cytomagalovirus; moloney murine leukemia virus; MoMuLV; chimeric;  
 KW alpha-lactalbumin; promoter; ds.

OS Chimeric - Encephalomyocarditis virus.

OS Chimeric - Bos Sp.

OS Chimeric - Cytomegalovirus.

OS Chimeric - Moloney murine leukemia virus.

OS Chimeric - Unidentified.

XX

FH Key Location/Qualifiers

FT promoter 1..812  
 FT /tag= a  
 FT /note= "CMV promoter/enhancer"  
 FT misc\_signal 853..855  
 FT /tag= b  
 FT /note= "MN14 antibody heavy chain gene signal peptide  
 FT start codon"  
 FT 2257..2259  
 FT /tag= c  
 FT /note= "MN14 antibody heavy chain gene start codon"  
 FT 2271..2846  
 FT /tag= d  
 FT /note= "EMCV IRES"  
 FT misc\_signal 2847..2849  
 FT /tag= e  
 FT /note= "Bovine alpha-lactalbumin signal peptide start  
 FT codon"  
 FT 2904..2906  
 FT /tag= f  
 FT /note= "First codon mature MN14 antibody light chain  
 FT gene"  
 FT 3543..3544  
 FT /tag= g  
 FT /note= "MN14 antibody heavy chain gene stop codon"  
 FT 3614..4207  
 FT /tag= h  
 FT /note= "MoMuLV 3' LTR"

WO200202783-A2.

10-JAN-2002.

29-JUN-2001; 2001WO-US20714.

03-JUL-2000; 2000US-215851P.

(GALA-) GALA DESIGN INC.

Bleek GT;

WPI; 2002-154749/20.

Novel regulatory elements including nucleic acid encoding hybrid  
 alpha-lactalbumin promoter or mutant RNA export element, for expressing  
 one or more proteins e.g. antibodies, pharmaceutical proteins in host  
 cells

Example 1; Fig 7; 151pp; English.

The invention relates to novel regulatory elements and vectors for the  
 expression of one or more proteins in a host cell. The invention further  
 provides methods of indirectly detecting the expression of a protein of  
 interest, comprising providing the host cell transfected with a vector  
 encoding a polycistronic sequence comprising a signal protein and a  
 desired protein operably linked by an internal ribosome entry site  
 (IRES), and culturing the host cell under suitable conditions so that  
 the signal protein and the desired protein is produced, where the  
 presence of signal protein indicates the presence of desired protein.  
 Regulatory elements and vectors of the invention are useful for the  
 expression of proteins of interest in a host cell. They are useful for  
 producing an immunoglobulin (Ig), preferably secretory Ig. They are  
 useful in the expression of one or more proteins such as erythropoietin,  
 growth hormone, insulin, immunoglobulins, protein C, cytokines and their  
 receptors, hormones, Von Willebrand's factor, lung surfactant, serum  
 albumins, DNase, vascular endothelial growth factor, receptors for  
 hormones or growth factors, rheumatoid factors, nerve growth factors,  
 CD proteins, osteoinductive factors, immunotoxins, bone morphogenetic  
 protein, interferons, colony stimulating factors, interleukins,  
 superoxide dismutase, T-cell receptor, surface membrane proteins,  
 viral antigens, transport proteins, addressins, regulatory proteins,  
 antibodies, chimeric proteins and their fragments. The vectors are  
 particularly useful for expressing G protein coupled receptors and other  
 transmembrane proteins. The retroviral vectors are useful for expressing



FT FT /\*tag= i  
XX XX /note= "MoMuLV 3' LTR"  
PN PN  
XX XX  
XX XX  
PD PD  
XX XX  
XX XX  
PF PF  
XX XX  
PR PR  
XX XX  
XX XX  
PA (GALA-) GALA DESIGN INC.  
XX XX  
XX XX  
PI Brémel RD, Miller LU, Bleck GT, York D;  
XX XX  
XX XX WPI; 2002-154737/20.  
DR WPI; 2002-154737/20.  
XX

WO200202738-A2.

10-JAN-2002.

29-JUN-2001; 2001WO-US20710.

03-JUL-2000; 2000US-215925P.

(GALA-) GALA DESIGN INC.

Brémel RD, Miller LU, Bleck GT, York D;

WPI; 2002-154737/20.

Host cell for producing a desired protein and for screening compounds useful for pharmaceutical, industrial, diagnostic and other purposes, comprises multiple integrating vectors having an exogenous gene -  
Example 1; Fig 11; 191pp; English.

The invention relates to a host cell comprising a genome having at least two integrated integrating vectors. The integrating vectors comprise at least one exogenous gene operably linked to a promoter. The host cell is useful for producing a desired protein and for comparing protein functions. The host cells comprises a reporter gene which is from gene fluorescent protein, luciferase, beta-galactosidase and beta-lactamase, and the assaying step further comprises detecting a signal from the reporter gene. The desired protein includes proteins for pharmaceutical, industrial, diagnostic and other purposes. The host cells is useful for indirectly detecting the expression of a desired protein, comprising providing the host cell transfected with a vector encoding a polyclonistic sequence comprising a signal protein and a desired protein operably linked by an internal ribosome entry site (IRES), and culturing the host cell under suitable conditions so that the signal protein and the desired protein is produced, where the presence of signal protein indicates the presence of desired protein. The present sequence is alpha lactalbumin (LA) Bot vector used in the invention. The vector comprises the following elements: bovine/human alpha-lactalbumin hybrid promoter, double mutated pre-mRNA processing enhancer (PPE) sequence; cc49 signal peptide, botulinum toxin antibody light chain, IRES from encephalomyocarditis virus (ECMV), bovine alpha-lactalbumin signal peptide, botulinum toxin antibody heavy chain, woodchuck mRNA processing enhancer (WPRE) sequence and 3' moloney murine leukemia virus (MoMuLV) LTR.

Sequence 3671 BP; 907 A; 868 C; 925 G; 971 T; 0 other;

Query Match 75.1%; Score 202.8; DB 24; Length 3671;  
Best Local Similarity 95.6%; Pred. No. 1e-55;  
Matches 262; Conservative 0; Mismatches 7; Indels 5; Gaps 5;

QY 1 GCTAGCTTAAGTAAAGCCGATTTTGCAGGCGATGGGAAATACATACTAGAGATAGAGA 60

DB 3108 GCTAGCTTAAGTAAAGCCGATTTTGCAGGCGATGGGAAATACATACTAGAGATAGAGA 3166

QY 61 AGTTTCAGATCAAGGTCAGGAACAGAGA-GAAACAGGAGATATGGGCCAAACAGGATATCTG 119

DB 3167 AGTTTCAGATCAAGGTCAGGAACAGAGATGGAGCTGAATATGGGCCAAACAGGATATCTG 3226

QY 120 TGGTAAAGCATGTTCTGCCCC-GCTCAGGGCCAGAACAGTGGAAACAGGAGAT-TGGGC 177

DB 3227 TGGTAAAGCATGTTCTGCCCCGCTCAGGGCCAGAACAGATGGAGCTGAATATGGGC 3286

QY 178 CAACAGGATATCTGTGGTAAGCAGTTCTGCCCC-GCTCAGGGCCAGAACAGATGGTTC 236

DB 3287 CAACAGGATATCTGTGGTAAGCAGTTCTGCCCCGCTCAGGGCCAGAACAGATGGTTC 3246

QY 237 CCAGATGCGGTCGCCGCCCTCAGCAGTTTCTAGA 270

DB 3347 CCAGATGCGGTCGCCGCCCTCAGCAGTTTCTAGA 3380

RESULT 14  
AAD28267

ID AAD28267 standard: DNA; 4207 BP.

XX AAD28267;

XX 22-APR-2002 (first entry)

XX Cytomegalovirus (CMV) MN14 vector.

XX Bovine: alpha-lactalbumin; promoter; pharmaceutical; industrial; ECMV;  
KW encephalomyocarditis virus; diagnostic; internal ribosome entry site;  
KW IRES; screening; CMV; cytomegalovirus; moloney murine leukemia virus;  
XX MoMuLV; chimeric; ds.

XX Chimeric - Encephalomyocarditis virus.

OS Chimeric - Bos sp.

OS Chimeric - Cytomegalovirus.

OS Chimeric - Moloney murine leukemia virus.

XX Chimeric - Unidentified.

XX Key Location/Qualifiers

FT Promoter 1..812

FT /\*tag= a

FT /note= "CMV promoter/enhancer"

FT misc\_signal 853..855

FT /\*tag= b

FT /note= "MN14 antibody heavy chain gene signal peptide start codon"

FT misc\_signal 2257..2259

FT /\*tag= c

FT /note= "MN14 antibody heavy chain gene start codon"

FT misc\_feature 2271..2846

FT /\*tag= d

FT /note= "EMCV IRES"

FT misc\_signal 2847..2849

FT /\*tag= e

FT /note= "Bovine alpha-lactalbumin signal peptide start codon"

FT misc\_signal 2904..2906

FT /\*tag= f

FT /note= "First codon mature MN14 antibody light chain gene"

FT misc\_signal 3543..3544

FT /\*tag= g

FT /note= "MN14 antibody light chain gene stop codon"

FT LTR 3614..4207

FT /\*tag= h

FT /note= "MoMuLV 3' LTR"

XX WO200202738-A2.

XX 10-JAN-2002.

XX 29-JUN-2001; 2001WO-US20710.

XX 03-JUL-2000; 2000US-215925P.

XX (GALA-) GALA DESIGN INC.

XX Brémel RD, Miller LU, Bleck GT, York D;

XX WPI; 2002-154737/20.

XX Host cell for producing a desired protein and for screening compounds useful for pharmaceutical, industrial, diagnostic and other purposes, comprises multiple integrating vectors having an exogenous gene -  
XX Example 1; Fig 7; 191pp; English.  
XX The invention relates to a host cell comprising a genome having at least

QY 237 CCAGATGGGTCCGCGCCCTCAGCAGTTTCTAGA 270  
 Db 270 CCCAGTGGGTCCAGCCCTCAGCAGTTTCTAGA 303

RESULT 12

AAF83096  
 ID AAF83096 standard; DNA; 3097 BP.

AC AAF83096;  
 XX 29-JUN-2001 (first entry)  
 DT Nucleotide sequence of a LTR plasmid.

XX Retrovirus; recombinase recognition sequence; RRS; LTR; recombinase;  
 KW long terminal repeat; pharmaceutical; cytosolic; antiinflammatory;  
 KW antirheumatic; antiarthritic; antiasthmatic; osteopathic; cardiant; MLV;  
 KW vasotropic; neuroprotective; nontropic; cerebroprotective; antipsoriatic;  
 KW antiarteriosclerotic; vulnerary; anti-HIV; antiulcer; thrombolytic;  
 KW dermatological; gene therapy; ss.

XX Synthetic.

WO200125466-A1.

PD 12-APR-2001.

PF 05-OCT-2000; 2000WO-GB03837.

PR 05-OCT-1999; 99GB-0023558.

PA (OXFO-) OXFORD BIOMEDICA UK LTD.

XX Slingsby J, Kingsman SM, Rohll J, Slade A;

PI WPI; 2001-281732/29.

XX Modifying producer cells for making retrovirus by transfecting with a  
 PT construct comprising a 5'- recombinase recognition sequence, long  
 PT terminal repeat and 3'- recombinase recognition sequence, in presence  
 PT of recombinase .

PS Example 5; Page 129-130; 133pp; English.

XX The invention relates to a method of modifying producer cells for making  
 CC retrovirus by transfecting with a construct comprising a 5'- recombinase  
 CC recognition sequence (RRS), long terminal repeat (LTR) and 3'- RRS, in  
 CC presence of recombinase. The regulated retroviral vector produced is  
 CC useful in the manufacture of a pharmaceutical composition to deliver a  
 CC NOI to a target site, and in the manufacture of a medicament for  
 CC diagnostic, therapeutic and/or medical applications. The recombinase  
 CC assisted method is useful for introducing regulated 3'-LTR into a derived  
 CC producer cell line to produce a high titer regulated retroviral vector.  
 CC The vector is useful in gene therapy for treating diseases like cancers,  
 CC inflammatory diseases, immunological disorders such as graft vs host  
 CC diseases, autoimmune diseases such as rheumatoid arthritis, allergic  
 CC diseases such as asthma, osteoporosis, cardiovascular diseases such as  
 CC congestive heart failure and ischemic heart disease, neurodegenerative  
 CC disorders such as multiple sclerosis, Alzheimer's disease, stroke and  
 CC cerebral ischemia, atherosclerosis, thrombotic disorders, dermatological  
 CC disorders such as atopic dermatitis, contact dermatitis and psoriasis,  
 CC wound healing, restenosis, infectious disorders such as HIV infections,  
 CC ulcers, digestive disorders such as anorexia, bulimia and cachexia, and  
 CC other diseases. The present sequence represents the nucleotide sequence  
 CC of a LTR plasmid.

SQ Sequence 3097 BP; 770 A; 823 C; 763 G; 741 T; 0 other;

Query Match 75.1%; Score 202.8; DB 22; Length 3097;  
 Best Local Similarity 95.6%; Pred. No. 9.7e-56;  
 Matches 262; Conservative 0; Mismatches 7; Indels 5; Gaps 5;

QY 1 GCTAGCTTAAGTAACGCCATTTTGCAGGCGATGGGAAAAATACATAACTGAGATAGAGA 60  
 Db 1 GCTAGCTTAAGTAACGCCATTTTGCAGGCGAT -GGAAANAATACATAACTGAGATAGAGA 59  
 QY 61 AGTTAGATCAAGGTCAGGAACAGA-GAAACAGAGAGAAATATGGCCCAACAGGATATCTG 119  
 Db 60 AGTTAGATCAAGGTCAGGAACAGAGTGAATATGGCCCAACAGGATATCTG 119  
 QY 120 TGGTAAGCAGTTCCTGCCCC-GCTCAGGCGCAAGAACAGTTGGAACAGGAGAAT-TGGGC 177  
 Db 120 TGGTAAGCAGTTCCTGCCCGCTCAGGCGCAAGAACAGATGGAACAGCTGAATATGGGC 179  
 QY 178 CAAACAGGATATCTGTGTAAGCAGTTCCTGCCCC-GCTCAGGCGCAAGAACAGATGCTC 236  
 Db 180 CAAACAGGATATCTGTGTAAGCAGTTCCTGCCCGCTCAGGCGCAAGAACAGATGCTC 239  
 QY 237 CCAGATGGGTCCGCGCCCTCAGCAGTTTCTAGA 270  
 Db 240 CCCAGATGGGTCCAGCCCTCAGCAGTTTCTAGA 273

RESULT 13

AAD28271

ID AAD28271 standard; DNA; 3671 BP.

AC AAD28271;

XX 22-APR-2002 (first entry)

DE Alpha-lactalbumin (LA) Bot vector.

XX Bovine; alpha-lactalbumin; promoter; pharmaceutical; industrial; BCMV;  
 KW encephalomyocarditis virus; diagnostic; internal ribosome entry site;  
 KW IRES; screening; CMV; cytomegalovirus; moloney murine leukemia virus;  
 KW MoMuLV; human; pre-mRNA processing enhancer; PPE; chimeric; WPRE;  
 KW woodchuck mRNA processing enhancer; botulinum; ds.

XX Chimeric - Encephalomyocarditis virus.

OS Chimeric - Bos sp.

OS Chimeric - Homo sapiens.

OS Chimeric - Moloney murine leukemia virus.

OS Chimeric - Marmota monax.

OS Chimeric - Clostridium botulinum.

OS Chimeric - Unidentified.

XX Key Location/Qualifiers

FT misc\_feature 1..2053

FT /\*tag= a

FT /note= "Bovine/human alpha-lactalbumin 5' flanking

FT region"

FT misc\_feature 2093..2336

FT /\*tag= b

FT /note= "Double mutated PPE sequence"

FT CDS 2387..2443

FT /\*tag= c

FT /product= "cc49 signal peptide coding region"

FT CDS 2444..3088

FT /\*tag= d

FT /product= "Bot antibody light chain Fab coding region"

FT misc\_feature 3112..3686

FT /\*tag= e

FT /note= "EMCV IRES"

FT CDS 3687..3745

FT /\*tag= f

FT /product= "Bovine alpha-lactalbumin signal peptide

FT coding region"

FT CDS 3746..4443

FT /\*tag= g

FT /product= "Bot antibody heavy chain Fab coding region"

FT misc\_feature 4481..5072

FT /\*tag= h

FT /note= "WPPE sequence"

FT LTR 5118..5711

Db 298 TGGTAAGCAGTTCTGCGCCGCTCAGGGCCAGGAACAGATGGAACAGCTGAATATGGGC 357  
QY 178 CAACAGGATATCTGTGTAAGCAGTTCTTCTGCCCC-GCTCAGGGCCAGGAACAGATGGTC 236  
Db 358 CAACAGGATATCTGTGTAAGCAGTTCTTCTGCCCGGCTCAGGGCCAGGAACAGATGGTC 417  
QY 237 CCCAGATGCGTCCCGCCCTCAGCAGTTCTTAGA 270  
Db 418 CCCAGATGCGTCCAGCCCTCAGCAGTTCTTAGA 451

RESULT 10  
AAQ54677  
ID AAQ54677 standard; DNA; 636 BP.  
XX  
AC AAQ54677;  
Y  
X 05-JUL-1994 (first entry)  
XX Promoter/Enhancer.  
XX  
DE  
XX  
KW Expression vector; promoter; enhancer; terminus; DMA1; DMA2;  
KW mutant; Namalwa cell; expression plasmid; repeat terminal;  
KW Moloney retrovirus; ss.  
XX  
XX Mus musculus.  
XX  
XX JP05317077-A.  
XX  
XX 03-DEC-1993.  
XX  
XX 25-MAY-1992; 92JP-0132650.  
XX  
XX 25-MAY-1992; 92JP-0132650.  
XX  
XX (KYOW ) KYOWA HAKKO KOGYO KK.  
XX  
XX WPI; 1994-011039/02.  
XX  
XX  
XX Prepn. of protein by recombinant animal cell - using expression  
PT vector with promoter or enhancer prep. from repeat terminal of  
PT Moloney retrovirus, and DNA for protein sequence, in Namalwa cell  
XX  
XX Claim 3; page 18; 34pp; Japanese.  
XX  
XX The sequence shows a promoter / enhancer DNA. The sequence can be  
XX recombined with a coding DNA within an expression plasmid which can  
XX be transformed into a Namalwa cell. This method can be used to prepare  
CC a useful protein efficiently.  
XX  
XX  
SQ Sequence 636 BP; 185 A; 160 C; 152 G; 139 T; 0 other;

Query Match 75.1%; Score 202.8; DB 15; Length 636;  
Best Local Similarity 95.6%; Pred. No. 4.7e-56;  
Matches 262; Conservative 0; Mismatches 7; Indels 5; Gaps 5;  
QY 1 GCTAGCTTAAGTAAGCCGCTTTTCAAGCGCATGGGAAAATACATAACTCAGAAATAGAGA 60  
Db 181 GCTAGCTTAAGTAAGCCGCTTTTCAAGCGCAT-GGAAAATACATAACTCAGAAATAGAGA 239  
QY 61 AGTTTCAGATCAAGGTTCAGGAACAGAGA-GAAACAGGAGAAATATGGCCCAACAGGATATCTG 119  
Db 240 AGTTTCAGATCAAGGTTCAGGAACAGAGTGGACAGCTGAATATGGCCCAACAGGATATCTG 299  
QY 120 TGGTAAGCAGTTCTTCTGCCCC-GCTCAGGGCCAGGAACAGATGGGAACAGGAGAAT-TGGGC 177  
Db 300 TGGTAAGCAGTTCTTCTGCCCGGCTCAGGGCCAGGAACAGATGGGAACAGGATATGGGC 359  
QY 178 CAACAGGATATCTGTGTAAGCAGTTCTTCTGCCCC-GCTCAGGGCCAGGAACAGATGGTC 236  
Db 360 CAACAGGATATCTGTGTAAGCAGTTCTTCTGCCCGGCTCAGGGCCAGGAACAGATGGTC 419  
QY 237 CCCAGATGCGGTCGCCCGCTCAGCAGTTTCTTAGA 270

Db 420 CCCAGATGCGGTCGCCCGCTCAGCAGTTTCTTAGA 453  
RESULT 11  
AAT32394  
ID AAT32394 standard; DNA; 702 BP.  
XX  
AC AAT32394;  
XX  
DT 28-SEP-1996 (first entry)  
XX  
DE Recombinant CMV/TAR MoMLV LTR in pMT-cat.  
XX  
KW MoMLV; long terminal repeat; CMV; HIV-1; tat protein; AIDS; retrovirus;  
KW transactivation response element; TAR; vector; antiviral; virucide;  
KW antitumour; tumour; cancer; gene therapy; ds.  
XX  
XX Chimeric Moloney murine leukemia virus;  
OS Chimeric human cytomegalovirus;  
OS Chimeric human immunodeficiency virus type 1.  
PN WO9614332-A1.  
XX  
PD 17-MAY-1996.  
XX  
XX 08-NOV-1995; 95WO-US14576.  
XX  
PR 08-NOV-1994; 94US-0336132.  
XX  
XX (CHAN/) CHANG L.  
XX  
XX Chang L;  
XX  
XX WPI; 1996-251713/25.  
XX  
XX Recombinant Moloney murine leukaemia long terminal repeat which is  
PT activated by HIV-1 Tat protein - useful in vectors for anti-viral,  
PT anti-tumour and gene therapy applications.  
XX  
XX Example 3; Page 86; 126pp; English.  
XX  
XX A recombinant MoMLV long terminal repeat (LTR) (AAT32394) was obtd.  
CC by replacing a portion of the U3 region of the Moloney murine  
CC leukaemia virus (MoMLV) LTR with the HIV-1 transactivation response  
CC element (TAR). Transfection studies showed the recombinant LTR  
CC to have a low basal activity and not to be responsive to Tat  
CC protein. This was in contrast to another recombinant LTR  
CC (AAT32382) that included the human cytomegalovirus immediate-early  
CC enhancer/promoter in addition to the TAR element.  
XX  
XX Sequence 702 BP; 170 A; 181 C; 193 G; 158 T; 0 other;

Query Match 75.1%; Score 202.8; DB 17; Length 702;  
Best Local Similarity 95.8%; Pred. No. 4.9e-56;  
Matches 262; Conservative 0; Mismatches 7; Indels 5; Gaps 5;  
QY 1 GCTAGCTTAAGTAAGCCGCTTTTCAAGCGCATGGGAAAATACATAACTGAGAATAGAGA 60  
Db 31 GCTAGCTTAAGTAAGCCGCTTTTCAAGCGCAT-GGAAAATACATAACTGAGAATAGAGA 89  
QY 61 AGTTTCAGATCAAGGTTCAGGAACAGAGA-GAAACAGGAGAAATATGGCCCAACAGGATATCTG 119  
Db 90 AGTTTCAGATCAAGGTTCAGGAACAGAGTGGACAGCTGAATATGGCCCAACAGGATATCTG 149  
QY 120 TGGTAAGCAGTTCTTCTGCCCC-GCTCAGGGCCAGGAACAGATGGGAACAGGAGAAT-TGGGC 177  
Db 150 TGGTAAGCAGTTCTTCTGCCCGGCTCAGGGCCAGGAACAGATGGGAACAGCTGAATATGGGC 209  
QY 178 CAACAGGATATCTGTGTAAGCAGTTCTTCTGCCCC-GCTCAGGGCCAGGAACAGATGGTC 236  
Db 210 CAACAGGATATCTGTGTAAGCAGTTCTTCTGCCCGGCTCAGGGCCAGGAACAGATGGTC 269



PI Von Laer M;  
 XX WPI; 2001-367622/38.  
 DR P-PSDB; AAB86198.  
 XX  
 PT New nucleic acid encoding membrane-anchored gp41 fusion protein, useful  
 PT for gene therapy of human immunodeficiency virus (HIV) infection,  
 PT prevents entry of virus into cells  
 XX  
 XX Claim 11; Page 28-30; 39pp; German.  
 PS  
 CC This invention describes a novel nucleic acid (I) comprising elements  
 CC that encode a signal peptide (SP) that provides transfer of expressed  
 CC polypeptide into the endoplasmic reticulum, a fragment (FI) of HIV gp41  
 CC protein, containing a segment from a heptad repeat region, a  
 CC transmembrane anchor (MSD) of a type I membrane protein and a flexible  
 CC linker (hinge) linking FI and MSD, therefore the formula of (I) is  
 CC : SP-FI-hinge-MSD. The products of the invention have antiviral activity  
 CC and act as HIV replication inhibitors. Vectors containing (I), also T  
 CC lymphocytes or hematopoietic stem cells transfected in vitro with (I),  
 CC are used in gene therapy of HIV infection.  
 XX  
 SQ Sequence 4148 BP; 880 A; 1177 C; 1103 G; 988 T; 0 other;  
 Query Match 77.1%; Score 208.2; DB 22; Length 4148;  
 Best Local Similarity 97.1%; Pred. No. 1.9e-57;  
 Matches 265; Conservative 0; Mismatches 3; Indels 5; Gaps 5;  
 QY 1 GCTAGCTTAAGTAAAGCCGATTTTCAAGGCGATGGGAAATACATACTGAGAATAGAGA 60  
 DB 3586 GCTAGCTTAAGTAAAGCCGATTTTCAAGGCGATGGGAAATACATACTGAGAATAGAGA 3643  
 QY 61 AGTTCAGATCAAGGTAGGAAACAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 120  
 DB 3644 AGTTCAGATCAAGGTAGGAAACAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 3703  
 QY 121 GGTAGGAGTTCCTCCGCC-CCTCAGGCGCCAGAGACAGTGGGAAACAGAGAGAT-TGGGCC 178  
 DB 3704 GGTAGGAGTTCCTCCGCCCGGCTCAGGCGCCAGAGACAGTGGGAAACAGAGAGAT-TGGGCC 3763  
 QY 179 AAACAGGATATCTGTGGTAAGCAGTTCCTCGCCCGGCTCAGGCGCCAGAGACAGATGGTCC 237  
 DB 3764 AAACAGGATATCTGTGGTAAGCAGTTCCTCGCCCGGCTCAGGCGCCAGAGACAGATGGTCC 3823  
 QY 238 CCAGATGGGTCCCGCCCTCAGCAGTTCCTCTAGA 270  
 O 3824 CCAGATGGGTCCCGCCCTCAGCAGTTCCTCTAGA 3856  
 RESULT 7  
 AAF83043  
 ID AAF83043 standard; DNA; 508 BP.  
 XX  
 AC AAF83043;  
 XX  
 DT 29-JUN-2001 (first entry)  
 DE  
 XX PCR amplified MLV U3 region from pHIT11.  
 XX  
 KW Retrovirus; recombinase recognition sequence; RRS; LTR; recombinase;  
 KW long terminal repeat; pharmaceutical; cytostatic; antiinflammatory;  
 KW antirheumatic; antiarthritic; antiasthmatic; osteopathic; cardiant; MLV;  
 KW vasotropic; neuroprotective; nootropic; cerebroprotective; antipsoriatic;  
 KW antiarteriosclerotic; vulnerary; anti-HIV; antiulcer; thrombolytic;  
 KW dermatological; gene therapy; ss.  
 XX  
 OS Murine leukemia virus.  
 XX  
 PN WO200125466-A1.  
 XX  
 PD 12-APR-2001.  
 XX  
 PF 05-OCT-2000; 2000WO-GB03837.

XX  
 PR 05-OCT-1999; 99GB-0023558.  
 XX  
 PA (OXFO-) OXFORD BIOMEDICA UK LTD.  
 XX  
 PI Slingsby J, Kingsman SM, Rohll J, Slade A;  
 DR WPI; 2001-281732/29.  
 XX  
 PT Modifying producer cells for making retrovirus by transfecting with a  
 PT construct comprising a 5'- recombinase recognition sequence, long  
 PT terminal repeat and 3'- recombinase recognition sequence, in presence  
 PT of recombinase  
 XX  
 PS Example 7; Page 51; 133pp; English.  
 XX  
 CC The invention relates to a method of modifying producer cells for making  
 CC retrovirus by transfecting with a construct comprising a 5'- recombinase  
 CC recognition sequence (RRS), long terminal repeat (LTR) and 3'- RRS, in  
 CC presence of recombinase. The regulated retroviral vector produced is  
 CC useful in the manufacture of a pharmaceutical composition to deliver a  
 CC NOI to a target site, and in the manufacture of a medicament for  
 CC diagnostic, therapeutic and/or medical applications. The recombinase  
 CC assisted method is useful for introducing regulated 3'-LTR into a derived  
 CC producer cell line to produce a high titer regulated retroviral vector.  
 CC The vector is useful in gene therapy for treating diseases like cancers,  
 CC inflammatory diseases, immunological disorders such as graft vs host  
 CC disease, autoimmune diseases such as rheumatoid arthritis, allergic  
 CC diseases such as asthma, osteoporosis, cardiovascular diseases such as  
 CC congestive heart failure and ischemic heart disease, neurodegenerative  
 CC disorders such as multiple sclerosis, Alzheimer's disease, stroke and  
 CC cerebral ischemia, atherosclerosis, thrombotic disorders, dermatological  
 CC disorders such as atopic dermatitis, contact dermatitis and psoriasis,  
 CC wound healing, restenosis, infectious disorders such as HIV infections,  
 CC ulcers, digestive disorders such as anorexia, bulimia and cachexia, and  
 CC other diseases. The present sequence represents a PCR amplified murine  
 CC leukemia virus (MLV) U3 region from pHIT11, used to replace Equine  
 CC infectious anemia virus (EIAV) PPT/U3 sequence. This is used in the  
 CC construction of EIAV vectors with LTR driven open reading frames.  
 XX  
 SQ Sequence 508 BP; 135 A; 129 C; 129 G; 115 T; 0 other;  
 Query Match 75.1%; Score 202.8; DB 22; Length 508;  
 Best Local Similarity 95.6%; Pred. No. 4.2e-56;  
 Matches 262; Conservative 0; Mismatches 7; Indels 5; Gaps 5;  
 QY 1 GCTAGCTTAAGTAAAGCCGATTTTCAAGGCGATGGGAAATACATACTGAGAATAGAGA 60  
 DB 58 GCTAGCTTAAGTAAAGCCGATTTTCAAGGCGATGGGAAATACATACTGAGAATAGAGA 116  
 QY 61 AGTTCAGATCAAGGTAGGAAACAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 119  
 DB 117 AGTTCAGATCAAGGTAGGAAACAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 176  
 QY 120 TGTAGCAGAGTTCCTCGCCCGGCTCAGGCGCCAGAGACAGTGGGAAACAGAGAT-TGGGCC 177  
 DB 177 TGTAGCAGAGTTCCTCGCCCGGCTCAGGCGCCAGAGACAGATGGGAAACAGAT-TGGGCC 236  
 QY 178 CAACAGGATATCTGTGGTAAGCAGTTCCTCGCCCGGCTCAGGCGCCAGAGACAGATGGTC 236  
 DB 237 CAACAGGATATCTGTGGTAAGCAGTTCCTCGCCCGGCTCAGGCGCCAGAGACAGATGGTC 296  
 QY 237 CCCAGATGGGTCCCGCCCTCAGCAGTTCCTCTAGA 270  
 DB 297 CCCAGATGGGTCCCGCCCTCAGCAGTTCCTCTAGA 330  
 RESULT 8  
 AA292704  
 ID AA292704 standard; cDNA; 614 BP.  
 XX  
 AC AA292704;  
 XX

XX The invention relates to a viral vector system for preparing recombinant  
 CC adeno-associated virus (AAV) particles comprising: at least two plasmid  
 CC vectors (ABL58983 and ABL58984) that include the two inverted terminal  
 CC repeats (ITR) of AAV and additional sequences and plasmid vectors without  
 CC ITRs but containing the rep and cap genes of AAV required for replication  
 CC and packaging. The system is useful for producing recombinant AAV for  
 CC production of a wide range of therapeutic glycoproteins in eukaryotic  
 CC cells. The system provides efficient, large scale production of  
 CC heterologous proteins in mammalian cells, without requiring an adenovirus  
 CC helper. It is not toxic to host cells and does not cause lysis, so  
 CC produced proteins are highly pure. The present sequence is that of the  
 CC pAIM E1B5K plasmid of the invention.  
 XX  
 SQ Sequence 6575 BP; 1623 A; 1566 C; 1695 G; 1691 T; 0 other;

Query Match 91.1%; Score 246; DB 24; Length 6575;  
 Best Local Similarity 99.3%; Pred. No. 1.le-69;  
 Matches 268; Conservative 0; Mismatches 0; Indels 2; Gaps 2;  
 QY 1 GCTAGCTTAAGTAAAGCCATTTTGAAGGCATGGGAAATAACATAACTCAGAAATAGAGA 60  
 432 GCTAGCTTAAGTAAAGCCCA-TTTCAGGCAAT-GGAAATAACATAACTCAGAAATAGAGA 489  
 61 AGTTCAGATCAAGGTTCAGGACAGAGAAACAGGAGAAATATGGGCCAAACAGGATATCTGT 120  
 490 AGTTCAGATCAAGGTTCAGGACAGAGAAACAGGAGAAATATGGGCCAAACAGGATATCTGT 549  
 121 GGTAAAGCATCTCTGCCCGCTCAGGGGCAAGAACAGAGTTGGACAGAGAAATGGGCCAA 180  
 550 GGTAAAGCATCTCTGCCCGCTCAGGGGCAAGAACAGAGTTGGACAGAGAAATGGGCCAA 609  
 181 ACAGGATATCTGTGTAAGCAGTTCTGCGCCGCTCAGGGGCAAGAACAGAGTTGGACAG 240  
 610 ACAGGATATCTGTGTAAGCAGTTCTGCGCCGCTCAGGGGCAAGAACAGAGTTGGACAG 669  
 241 GATCGGTCGCCCGCTCAGCAGTTCTCTAGA 270  
 670 GATCGGTCGCCCGCTCAGCAGTTCTCTAGA 699

RESULT 5  
 AAA96220 standard; DNA; 9830 BP.  
 ID AAA96220  
 AC AAA96220;  
 DT 08-FEB-2001 (first entry)  
 XX Nucleotide sequence of pTRONIN, a retroviral vector.  
 DE Retroviral vector; inflammatory disorder; dermatological disorder;  
 XX cardiovascular disorder; autoimmune disease; neurological disorder;  
 XX cancer; gene therapy; ss.  
 XX Synthetic.  
 XX WO200056910-A1.  
 XX 28-SEP-2000.  
 XX 22-MAR-2000; 2000WO-GB01091.  
 XX 22-MAR-1999; 99GB-0006615.  
 XX (OXFO-) OXFORD BIOMEDICA UK LTD.  
 XX Uden M, Mitrophanous K;  
 XX WPI; 2000-628271/60.  
 XX Retroviral vector for delivering one or more nucleotides of interest to  
 PT a target site has functional and non-functional splice donor and splice

PT acceptor sites -  
 XX  
 PS Example 8; Fig 33; 148pp; English.  
 XX  
 CC The specification describes a retroviral vector which comprises a  
 CC nucleotide sequence of interest flanked by functional splice donor  
 CC sites and functional splice acceptor sites. The vector is derived from  
 CC a retroviral pro-vector. The retroviral vector is useful for preparing  
 CC pharmaceutical compositions to deliver one or more nucleotide sequences  
 CC of interest to a target site. The retroviral vectors are especially  
 CC useful for treating inflammatory disorders, cancers, dermatological  
 CC disorders, cardiovascular disorders, autoimmune diseases and neurological  
 CC disorders. The retroviral vector is useful in gene therapy. The present  
 CC sequence represents the retroviral vector pTRONIN, a vector of the  
 CC invention.  
 XX  
 SQ Sequence 9830 BP; 2209 A; 2659 C; 2688 G; 2274 T; 0 other;  
 Query Match 79.7%; Score 215.2; DB 21; Length 9830;  
 Best Local Similarity 94.1%; Pred. No. 1.5e-59;  
 Matches 256; Conservative 0; Mismatches 13; Indels 3; Gaps 3;  
 QY 1 GCTAGCTTAAGTAAAGCCATTTTGAAGGCATGGGAAATAACATAACTCAGAAATAGAGA 60  
 6141 GCTAGCTTAAGTAAAGCCATTTTGAAGGCAT-GGAAATAACATAACTCAGAAATAGAGA 6199  
 61 AGTTCAGATCAAGGTTCAGGACAGAGAAACAGGAGAAATATGGGCCAAACAGGATATCTGT 119  
 6200 AGTTCAGATCAAGGTTCAGGACAGAGAAACAGGAGAAATATGGGCCAAACAGGATATCTGT 6259  
 120 TGGTAAAGCATCTCTGCGCCGCTCAGGGGCAAGAACAGAGTTGGACAGAGAAATGGGCC 178  
 6260 TGGTAAAGCATCTCTGCGCCGCTCAGGGGCAAGAACAGAGTTGGACAGAGAAATGGGCC 6319  
 179 AAACAGGATATCTGTGTAAGCAGTTCTGCGCCGCTCAGGGGCAAGAACAGAGTTGGTCCC 238  
 6320 AAACAGGATATCTGTGTAAGCAGTTCTGCGCCGCTCAGGGGCAAGAACAGAGTTGGTCCC 6379  
 239 CAGATGCGGTCCCGCTCAGCAGTTCTCTAGA 270  
 6380 CAGATGCGGTCCCGCTCAGCAGTTCTCTAGA 6411

RESULT 6  
 AAH20890 standard; DNA; 4148 BP.  
 ID AAH20890  
 AC AAH20890;  
 DT 24-AUG-2001 (first entry)  
 XX Vector containing HIV gp41 DNA SEQ ID 1.  
 DE Transmembrane anchor; gene therapy; endoplasmic reticulum; gp41;  
 XX antiviral; HIV replication inhibitor; T lymphocyte; viral infection;  
 KW hematopoietic stem cell; ds.  
 XX Human immunodeficiency virus type 1.  
 XX  
 XX Key Location/Qualifiers  
 FH CDS 1438..1773  
 FT /\*tag= a  
 FT  
 XX WO200137881-A2.  
 XX 31-MAY-2001.  
 XX 24-NOV-2000; 2000WO-EPI1733.  
 XX 25-NOV-1999; 99DE-1057838.  
 XX (PETT-) PETTE INST HEINRICH.  
 XX

CC at least two poliovirus derived internal ribosomal entry site (IRES)  
 CC sequences. The vector can be expressed in mammalian host cells for the  
 CC production of heteromeric fusion proteins. This expression system is  
 CC claimed to produce the heteromeric proteins in high yields.

XX Sequence 8298 BP; 1974 A; 2364 C; 2134 G; 1826 T; 0 other;  
 SQ Query Match 100.0%; Score 270; DB 19; Length 8298;  
 Best Local Similarity 100.0%; Pred. No. 1.8e-77;  
 Matches 270; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCTAGCTTAAGTACGCGCATTTGCAAGCGCATGGGAAATAATACATACTGAGAAATAGAGA 60  
 DB 36 GCTAGCTTAAGTACGCGCATTTGCAAGCGCATGGGAAATAATACATACTGAGAAATAGAGA 95  
 QY 61 AGTTCAGATCAAGGTTCAGGAAACAGAGAGAGAAATATGGCCCAACAGGATATCTGT 120  
 DB 96 AGTTCAGATCAAGGTTCAGGAAACAGAGAGAGAAATATGGCCCAACAGGATATCTGT 155  
 QY 121 GGTAAAGCAGTTCTCGCCCGCTCAGGCGCAAGAACAGAGTTGGGAGAAATGGGCCAA 180  
 DB 156 GGTAAAGCAGTTCTCGCCCGCTCAGGCGCAAGAACAGAGTTGGGAGAAATGGGCCAA 215  
 QY 181 ACAGGATATCTGTGTAAGCAGTTCTCGCCCGCTCAGGCGCAAGAACAGAGTTGGGAGAAATGGGCCAA 240  
 DB 216 ACAGGATATCTGTGTAAGCAGTTCTCGCCCGCTCAGGCGCAAGAACAGAGTTGGGAGAAATGGGCCAA 275  
 QY 241 GATCGGTCCCGCCCTCAGCAGTTCTTAGA 270  
 DB 276 GATCGGTCCCGCCCTCAGCAGTTCTTAGA 305

RESULT 3  
 AAV08560/c  
 ID AAV08560 standard; cDNA: 3263 BP.  
 XX AC AAV08560;  
 XX DT 12-FEB-1999 (first entry)

XX Transgene for epitope tagged TBP protein.  
 XX TATA-box binding protein; epitope-tagged TBP; transcription complex; TAF;  
 KW TBP associated factor; TAF-interaction factor; gene expression regulator;  
 KW ss.  
 X Homo sapiens.  
 S Synthetic.  
 OS EP881288-Al.  
 XX 02-DEC-1998.  
 XX 26-MAY-1998; 98EP-0109516.  
 XX 26-MAY-1997; 97EP-0108433.  
 XX (FARH ) HOECHST AG.  
 XX Berglund E, Kirschbaum B, Meisterernst M, Polites G;  
 XX WPI; 1999-001394/01.

XX Transgenic animal expressing epitope-tagged TATA-box binding protein  
 PT - for isolating higher-order transcription complexes and specific  
 PT factors that associate with the protein, useful as potential  
 PT therapeutic agents  
 XX Claim 22; Page 27-29; 38pp; English.  
 PS This sequence encodes an epitope-tagged TATA-box binding protein (TBP)  
 CC that is expressed by the transgenic non-human animals of the invention.  
 CC The animals are used to produce TBP. TBP is used to isolate and

CC characterise higher-order transcription complexes (from different tissue  
 CC and cell types, optionally at different developmental stages). It is also  
 CC used to identify new and/or specific TBP associated factors (TAFs,  
 CC e.g. transcription factors, activators or inhibitors) and TAF-interaction  
 CC factors, and to raise antibodies against TBP. The TAFs may be useful for  
 CC regulating gene expression, e.g. disease-related genes, so are potential  
 CC pharmaceuticals, also for identifying human analogues for use in drug  
 CC screening. The antibodies are used for affinity purification of TBP and  
 CC its complexes. TBP can isolate transcription complexes from a wide  
 CC variety of different tissues and cells (contrast known methods that are  
 CC limited to isolation from a particular cell type).

XX Sequence 3263 BP; 860 A; 794 C; 756 G; 853 T; 0 other;  
 SQ Query Match 94.3%; Score 254.6; DB 20; Length 3263;  
 Best Local Similarity 96.7%; Pred. No. 1.2e-72;  
 Matches 260; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

QY 1 GCTAGCTTAAGTACGCGCATTTTGCAGGCGCATGGGAAATAATACATACTGAGAAATAGAGA 60  
 DB 289 GCTAGCTTAAGTACGCGCATTTTGCAGGCGCATGGGAAATAATACATACTGAGAAATAGAGA 230  
 QY 61 AGTTCAGATCAAGGTTCAGGAAACAGAGAGAGAAATATGGCCCAACAGGATATCTGT 120  
 DB 229 AGTTCAGATCAAGGTTCAGGAAACAGAGAGAGAAATATGGCCCAACAGGATATCTGT 170  
 QY 121 GGTAAAGCAGTTCTCGCCCGCTCAGGCGCAAGAACAGAGTTGGGAGAAATGGGCCAA 180  
 DB 169 GGTAAAGCAGTTCTCGCCCGCTCAGGCGCAAGAACAGAGTTGGGAGAAATGGGCCAA 110  
 QY 181 ACAGGATATCTGTGTAAGCAGTTCTCGCCCGCTCAGGCGCAAGAACAGAGTTGGGAGAAATGGGCCAA 240  
 DB 109 ACAGGATATCTGTGTAAGCAGTTCTCGCCCGCTCAGGCGCAAGAACAGAGTTGGGAGAAATGGGCCAA 50  
 QY 241 GATCGGTCCCGCCCTCAGCAGTTCTTAG 269  
 DB 49 GATCGGTCCCGCCCTCAGCAGTTCTTAG 21

RESULT 4  
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 XX AC ABL58984;  
 XX DT 22-JUL-2002 (first entry)  
 XX AAV expression vector pAIM ELB55K SEQ ID NO 2.  
 DE Adeno-associated virus; AAV; glycoprotein; ITR; cap; rep; vector; ds.  
 XX Adeno-associated virus.  
 OS Synthetic.  
 OS WO200238782-A2.  
 XX 16-MAY-2002.  
 XX 13-NOV-2001; 2001WO-EPI3125.  
 XX 13-NOV-2000; 2000DE-1056210.  
 XX (ARIM-) ARIMEDES BIOTECHNOLOGY GMBH.  
 XX Orberger G, Hellmuth K, Wagener C;  
 XX WPI; 2002-435853/46.  
 PT Vector system for preparing recombinant adeno-associated viral  
 PT particles, used for high-level expression of heterologous therapeutic  
 PT proteins in eukaryotic cells -  
 XX Claim 43; Page 52-56; 59pp; German.

mammalian gene expression, is based on the interferon regulating factor-1 and its binding site

Claim 1; Page 9; 19pp; English.

The present invention describes a promoter-transactivator system for inducible high-level expression of mammalian genes, and optionally for control of cell growth. The promoter-transactivator system comprises: (i) a promoter construct (IRFE promoter); and (ii) transactivator construct encoding a fusion protein of IRF-1 (interferon regulating factor-1) and the estrogen receptor (ER). The IRFE promoter construct has the structure (MPSV-E)-(IRF-1 binding site)-(CMV)-DNA where MPSV-E indicates the myeloproliferative sarcoma virus enhancer repeat given in AAC64591; IRF-1 binding site given in AAC64592; and CMV is the cytomegalovirus minimal promoter given in AAC64593 or their functionally equivalent variants with one or more nucleotides substituted, inserted or deleted. The promoter transactivator system is a transcription regulator. Increased transcription results from binding of the IRFE promoter to the transactivator, which is activated by oestradiol or other ER ligands (these compounds displace the heat-shock protein 90 which normally binds to the IRF-1/ER fusion, preventing its activation). The system is used for increasing production of therapeutically active proteins and where a IRF-1-green fluorescent protein-human ER construct is used, for fluorescence-activated cell sorting/analysis of transfected cells.

Sequence 270 BP; 85 A; 61 C; 74 G; 50 T; 0 other;

Query Match 100.0%; Score 270; DB 21; Length 270;  
Best Local Similarity 100.0%; Pred. No. 3.7e-78;  
Matches 270; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 61 AGTTCAGATCAAGGTCAGAGAACAGAGAGAAATATGGGCCAAACAGGATATCTGT 120

Qy 121 GGTAGAGAGTTCTCCCGCGCTCAGGCGCAAGAACAGAGAGAAATGGGGCCAA 180  
Db 121 GGTAGAGAGTTCTCCCGCGCTCAGGCGCAAGAACAGAGAGAAATGGGGCCAA 180

Qy 181 ACAGGATATCTGTGTAGACAGTTCTGCGCGCTCAGGCGCAAGAACAGAGATGTCGCCA 240  
Db 181 ACAGGATATCTGTGTAGACAGTTCTGCGCGCTCAGGCGCAAGAACAGATGTCGCCA 240

Qy 241 GATCGGTCGCGCGCTCAGCAGTTTCTAGA 270  
Db 241 GATCGGTCGCGCGCTCAGCAGTTTCTAGA 270

RESULT 2  
AAV18096  
ID AAV18096 standard; DNA; 8298 BP.  
AC AAV18096;  
XX  
XX  
XX  
DT 04-AUG-1998 (first entry)  
XX  
XX  
DE pmCLDHPA triclitrone vector for the expression of hmAb45-TNF alpha.  
XX  
XX Circular; antibody-cytokine fusion protein; triclitrone vector;  
KW TNF alpha; IL-2; IRES; internal ribosome entry site; ds.  
XX  
XX Synthetic.  
OS  
XX Key Location/Qualifiers  
XX promoter 1..904  
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FT /note= "CMV promoter with an upstream MPSV enhancer"  
FT intron 905..976

FT /tag= b  
FT /number= Intron 1  
FT 977..1018  
FT /tag= c  
FT /note= "Partial leader sequence"  
FT 1019..1106  
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FT /note= "5'UTR from poliovirus"  
FT 1107..1433  
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FT /product= "Light chain hmAb425, variable region"  
FT 1107..1115  
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FT /note= "Rest of the leader sequence"  
FT 1434..1595  
FT /tag= g  
FT /number= Intron 2  
FT 1596..1913  
FT /tag= h  
FT /product= "Light chain hmAb425, constant region"  
FT 1914..2028  
FT /tag= i  
FT /note= "5' UTR from poliovirus"  
FT 2029..2159  
FT /tag= j  
FT /note= "Poliovirus derived internal ribosome entry site"  
FT 1260..2581  
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FT 2582..4537  
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FT /note= "Heavy chain hmAb425 fused to TNF alpha"  
FT 4565..5279  
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FT /note= "Comprises of a 5' UTR from poliovirus, an internal ribosome entry site and intron 4"  
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FT /tag= n  
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FT /note= "Selection marker"  
FT 5929..6181  
FT /tag= o  
FT /standard\_name= SV40 PolyA  
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XX WO9811241-A1.  
XX  
XX 19-MAR-1998.  
XX  
XX 02-SEP-1997; 97WO-EP04765.  
XX  
XX 30-SEP-1996; 96EP-0115635.  
XX  
XX 16-SEP-1996; 96EP-0114820.  
XX  
XX (MERE ) MERCK PATENT GMBH.  
XX  
XX Brummer W, Burge C, Dunker R, Hauser H, Mielke C;  
PI Rieke E, Von Hoegen I, Welge T;  
XX  
XX WPI; 1998-207400/18.  
XX  
XX P-PSDB; AAW48647, AAW48648, AAW48649, AAW48650, AAW48651.  
XX  
XX Oligo:clitrone expression vector - useful for production of, e.g.  
PT MAb425/TNF-a or MAb425/IL-2 antibody fusion protein  
XX  
XX Claim 11; Fig 15; 89pp; English.  
XX  
XX The present sequence represents a new pmCLDHPA triclitrone vector for  
CC the expression of an antibody-cytokine fusion protein, hmAb425-TNF  
CC alpha. hmAb425-TNF alpha comprises of the TNF alpha fused to the  
CC C-terminus of the heavy chain of humanized monoclonal antibody 425.  
CC The TNF alpha sequence can be substituted by the IL-2 sequence. The  
CC hmAb425 has specificity for the human EGF receptor. The vector also  
CC contains a strong promoter/enhancer unit, a selection marker gene and



GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: March 5, 2003, 22:29:12 ; Search time 263.478 Seconds  
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SUMMARIES

Result No.	Query Match	Score	Length	ID	Description
1	270	100.0	270	21 AAC64591	Myeloproliferative
2	270	100.0	8298	19 AAV18096	PMCLDHAP tricitro
3	254.6	94.3	3263	20 AAV08560	Transgene for epit
4	246	91.1	6575	24 ABL38984	AAV expression vec
5	215.2	79.7	9830	21 AAA96220	Nucleotide sequenc
6	208.2	77.1	4148	22 AAH20890	Vector containing
7	202.8	75.1	508	22 AAF3043	PCR amplified MLV
8	202.8	75.1	614	21 AAT92704	Moloney murine leu
9	202.8	75.1	635	18 AAT97160	Moloney long termi

10	202.8	75.1	636	15 AAQ54677	Promoter/Enhancer.
11	202.8	75.1	702	17 AAT32394	Recombinant CMV/TA
12	202.8	75.1	3097	22 AAF83096	Nucleotide sequenc
13	202.8	75.1	3671	24 AAD28271	Alpha-lactalbumin
14	202.8	75.1	4207	24 AAD28267	Cytomegalovirus (C
15	202.8	75.1	4207	24 AAD28306	Cytomegalovirus (C
16	202.8	75.1	4210	24 AAD28268	Cytomegalovirus (C
17	202.8	75.1	4210	24 AAD28307	Cytomegalovirus (C
18	202.8	75.1	4661	24 AAD28273	Alpha-lactalbumin
19	202.8	75.1	4661	24 AAD28312	Alpha-lactalbumin
20	202.8	75.1	4776	20 AAX77617	Expression constru
21	202.8	75.1	4776	20 AAX77614	Expression constru
22	202.8	75.1	4924	21 AAZ34937	Retrovirus vector.
23	202.8	75.1	4924	22 AAF30946	Retrovirus vector.
24	202.8	75.1	5109	18 AAT76800	PLJ Rev m10 retrov
25	202.8	75.1	5130	24 AAD28311	LSRNL vector. Chl
26	202.8	75.1	5176	18 AAT76801	PLJ mutant Rev m10
27	202.8	75.1	5177	20 AAZ05997	Bovine scavenger r
28	202.8	75.1	5594	19 AAV33629	GENSA 981, a monom
29	202.8	75.1	5617	24 AAD32077	Human alpha-1-anti
30	202.8	75.1	5689	20 AAZ11445	Retroviral vector
31	202.8	75.1	5689	20 AAX61061	Retroviral vector
32	202.8	75.1	5689	21 AAA96208	Nucleotide sequenc
33	202.8	75.1	5691	24 AAD28274	Alpha-lactalbumin
34	202.8	75.1	5691	24 AAD28313	Alpha-lactalbumin
35	202.8	75.1	5711	24 AAD28310	Alpha-lactalbumin
36	202.8	75.1	5715	21 AAZ34936	Retrovirus vector.
37	202.8	75.1	5715	22 AAF30945	Vector used in inv
38	202.8	75.1	5731	24 AAD32078	Human alpha-1-anti
39	202.8	75.1	5732	24 AAD28269	Mouse mammary tumo
40	202.8	75.1	5732	24 AAD28308	Mouse mammary tumo
41	202.8	75.1	5865	19 AAV04002	Retroviral vector
42	202.8	75.1	5874	20 AAX90484	Plasmid retroviral
43	202.8	75.1	6026	24 AAD32075	Human albumin prom
44	202.8	75.1	6140	24 AAD32076	Human albumin prom
45	202.8	75.1	6141	20 AAX90481	Plasmid retroviral

ALIGNMENTS

RESULT 1

AAC64591  
ID AAC64591 standard; DNA; 270 BP.

XX AAC64591;

XX AC

XX 15-FEB-2001 (first entry)

XX Myeloproliferative sarcoma virus enhancer sequence SEQ ID NO:1.

XX Myeloproliferative sarcoma virus; MPSV; MPSV-E; enhancer; CMV;

XX Cytomegalovirus; firefly; IRF-1 binding site; minimal promoter;

XX interferon regulatory factor 1 binding site; transcription regulator;

XX promoter transactivator system; ds.

XX Myeloproliferative sarcoma virus.

XX EPI046710-A1.

XX 25-OCT-2000.

XX 23-APR-1999; 99EP-0108068.

XX 23-APR-1999; 99EP-0108068.

XX (GBFB ) GES BIOTECHNOLOGISCHE FORSCHUNG MBH.

XX Mueller PP, Geserick C, Schroeder K, Hauser H;

XX WPI; 2000-648930/63.

XX Promoter-transactivator system, useful for inducing high level

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Best Local Similarity 94.1%; Pred. No. 1.6e-53;
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Qy 120 TGGTAAGCAGTTCTTGGCCCTC-GCTCAGGGCCAAAGAACAGTTGGAACAGGAAATTTGGGCC 178
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.y 238 CCAGATGCGGTCCTCGCCCTCAGCAGTTTCTAGA 270
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Job time : 1503.61 secs

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Qy 61 AGTTTCAGATCAGGTCAGGAACAGAGAGAGAGAGATATGGGCAACAGGATATCTGT 120  
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Db 3644 AGTTTCAGATCAGGTCAGGAACAGAGAGAGAGATATGGGCAACAGGATATCTGT 3703  
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3704 GCTAGCAGCTTCTGCCCCGGCTCAGGCGCAAGAACAGGTTGGACAGAGATATGGGCC 3763

Db 3704 GCTAGCAGCTTCTGCCCCGGCTCAGGCGCAAGAACAGGTTGGACAGAGATATGGGCC 3763

Qy 179 AAACAGGATATCTGTGTAAGCAGTCTCTGCCCC-GCTCAGGCGCAAGAACAGATGGTCC 237  
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3764 AAACAGGATATCTGTGTAAGCAGTCTCTGCCCCGGCTCAGGCGCAAGAACAGATGGTCC 3823

Qy 238 CCAGATGGCGTCCCGCCCTCAGCAGTTTCTAGA 270  
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RESULT 14  
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LOCUS

DEFINITION Sequence 6 from patent US 5858744.  
ACCESSION AR028672  
VERSION AR028672.1 GI:5940645  
KEYWORDS Unknown.  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 9318)  
AUTHORS Baum,C., Stocking-Harbers,C. and Osterlag,W.  
TITLE Retroviral vector hybrids and the use thereof for gene transfer  
JOURNAL Patent: US 5858744-A 6 12-JAN-1999;  
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5957 AGTTTCAGATCAGGTCAGGAACAGAGAGAGAGATATGGGCAACAGGATATCTGT 6016

Db 5957 AGTTTCAGATCAGGTCAGGAACAGAGAGAGAGATATGGGCAACAGGATATCTGT 6016

Qy 121 GCTAGCAGTCTCTGCCCC-GCTCAGGCGCAAGAACAGGTTGGACAGAGAGAT-TGGGCC 178  
|||||  
6017 GCTAGCAGTCTCTGCCCCGGCTCAGGCGCAAGAACAGGTTGGACAGAGATATGGGCC 6076

Db 6017 GCTAGCAGTCTCTGCCCCGGCTCAGGCGCAAGAACAGGTTGGACAGAGATATGGGCC 6076

Qy 179 AAACAGGATATCTGTGTAAGCAGTCTCTGCCCC-GCTCAGGCGCAAGAACAGATGGTCC 237  
|||||  
6077 AAACAGGATATCTGTGTAAGCAGTCTCTGCCCCGGCTCAGGCGCAAGAACAGATGGTCC 6136

Db 6077 AAACAGGATATCTGTGTAAGCAGTCTCTGCCCCGGCTCAGGCGCAAGAACAGATGGTCC 6136

Qy 238 CCAGATGGCGTCCCGCCCTCAGCAGTTTCTAGA 270  
|||||  
6137 CCAGATGGCGTCCCGCCCTCAGCAGTTTCTAGA 6169

Db 6137 CCAGATGGCGTCCCGCCCTCAGCAGTTTCTAGA 6169

RESULT 15  
MLM3LTR  
LOCUS  
DEFINITION Moloney murine leukemia virus TB 3' LTR region.  
ACCESSION M24204  
VERSION M24204.1 GI:600759  
KEYWORDS LTR.  
SOURCE Moloney murine leukemia virus (strain TB) RNA.  
ORGANISM Moloney murine leukemia virus  
VIRUSES; Retroviral viruses; Retroviridae; Gammaretrovirus.  
REFERENCE 1 (bases 1 to 733)  
AUTHORS Yuen,P.H. and Szurek,P.F.  
TITLE The reduced virulence of the thymotropic Moloney murine leukemia  
virus derivative MoMuLV-TB is mapped to 11 mutations within the U3  
region of the long terminal repeat  
J. Virol. 63 (2), 471-480 (1989)  
JOURNAL 89094971  
MEDLINE 2783465  
PUBMED  
COMMENT On Dec 10, 1994 this sequence version replaced gi:341122.  
FEATURES Location/Qualifiers  
source  
1..733  
/organism="Moloney murine leukemia virus"  
/strain="TB"  
/db\_xref="taxon:11801"  
repeat\_region 140..590  
/note="U3 region"  
repeat\_region 140..153  
/rpt\_type=inverted  
repeat\_region 245..319  
/rpt\_type=direct  
protein\_bind 246..251

FEATURES	Location/Qualifiers
source	1. 2854
	/organism="Myeloproliferative sarcoma virus"
	/db_xref="taxon:11813"
CDS	973..2001
	/note="v-mos protein"
	/codon_start=1
	/protein_id="AAA46580.1"
	/db_xref="GI:332209"
	/translation="MPSPLSLCYLPRELSPVDSRCSPLVAPRKAGKFLGTTGPP RAGLPRLRAFWISIDWEQVCLMRGLSGGGSVYKATYHGVPAIKQVKNCTKDLRAS QRSFALNIAIRLHDNINRYVAASRTPEDSNLGTIIMEFGVNTLHQVIYGA PEPSREQSLQSLKISUDVYVGLFHSQSLHLDLAPANILISEQDVCKISDFG CSQKLQRCRQSPHHIGGTYHOAPEILKGAITPRADIIYSGFIILQMWTREVYIS GEQYVQYAVVAYNLRLPSLTGAVETASLTGKTLQNIQNCWEARALQRPAGELLQRLD KAFRGALG"
LTR	2268..2854
	/note="3', long terminal repeat"
repeat_region	2383..2456
	/note="tandem repeat A"
repeat_region	2457..2530
	/note="tandem repeat B"
BASE COUNT	696 a 814 c 709 g 635 t
ORIGIN	1 bp upstream of HindIII site.
Query Match	82.6%; Score 223; DB 14; Length 2854;
Best Local Similarity	98.5%; Pred. No. 1.3e-58;
Matches 267; Conservative	0; Mismatches 0; Indels 4; Gaps 4;
Qy 1	GCTAGCTTAAGTACGCCATTTTCGAGGCAATGGGAAAATACTACTAGCATAGAGA 60
Db 2298	GCTAGCTTAAGTACGCCA-TTTTCGAGGCAT-GGAAAAATACATACTGAGATAGAGA 2355
Qy 61	AGTTCAGATCAAGCTCAGGAACAGGAACAGGAGAAATATGGCCAAACAGGATATCTGT 120
Db 2356	AGTTCAGATCAAGGTCAGGAACAGGAACAGGAGAAATATGGCCAAACAGGATATCTGT 2415
Qy 121	GGTAAGCAGTTCCTGCCCCGCTCAGGGCCAAAGAACACAGTTGGAACAGGAGAAT-TGGGCCA 179
Db 2416	GGTAAGCAGTTCCTGCCCCGCTCAGGGCCAAAGAACACAGTTGGAACAGGAGAATATGGGCCA 2475
Qy 180	AACAGGATATCTGTGTAAGCAGGTTCCTGCCCGCTCAGGGCCCAAGAACACAGTGGTCCCC 239
Db 2476	AACAGGATATCTGTGTAAGCAGGTTCCTGCCCGCTCAGGGCCCAAGAACACAGTGGTCCCC 2535
Y 240	AGATCGGTCGCCGCCCTCAGCAGGATTTCTAGA 270
Db 2536	AGAT-CGGTCCCGCCCTCAGCAGGATTTCTAGA 2565
RESULT 12	
S66424/c	
LOCUS	S66424 544 bp DNA linear ROD 06-JAN-1994
DEFINITION	c-myc [5' LTR, provirus insertion] [mice, plasmacytoma RFPC 2782, Genomic Mutant, 544 nt].
ACCESSION	S66424
VERSION	S66424.1
KEYWORDS	GI:439771
SOURCE	Mus sp. plasmacytoma RFPC 2782.
ORGANISM	Mus sp.
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 544)
AUTHORS	Shaugnessy J.D. Jr., Owens J.D. Jr., Wiener F., Hilbert D.M., Huppi K., Potter M. and Mushinski J.F.
TITLE	Retriable enhancer insertion 5' of c-myc in two translocation-negative mouse plasmacytomas upregulates c-myc expression to different extents
JOURNAL	Oncogene 8 (11), 3111-3121 (1993)
MEDLINE	94020837
PUBMED	8414513
REMARK	GenBank staff at the National Library of Medicine created this entry [NCBI qibbsq 138854] from the original journal article.

This sequence comes from Fig. 4b.

```

FEATURES             source          1..544
   Location/Qualifiers
     /organism="Mus sp."
     /db_xref="taxon:10095"
gene                 1..544
     /gene="c-myc"
BASE COUNT           121 a      129 c      117 g      177 t
ORIGIN

Query Match              77.4%   Score 209;   DB 10;   Length 544;
Best Local Similarity    94.9%;   Pred.No. 2.4e-54;
Matches 259; Conservative 0; Mismatches 10; Indels 4; Gaps 4;

Qy  1  GCTAGCTTAAGTAACGGCATTTCGAAGGCATGGGAAAAATACATAACTGAGAATAGAGA  60
      | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db  392 GCTACGTTAAGTAACGGCATTTCGAAGGCAT -GGAAAAATAGATAACTGAGAATAGAGA  334

Qy  61 AGTTCAGATCAAGGTCAGGAACACAGAGAACAAGGAGAGATATGGGCCAAACAGGATATCTGT  120
      | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db  333 AGTTCAGATCAAGGTCAGGAACACAGAGAACAAGTGAATATATGGGCCAAACAGGATATCTGT  274

Qy  121 GGTAAGCAGTTCCTGGCCCC -GCTCAGGCGCCAAAGAACAGTTCGGAACAGAGAGAT -TGGGCC  178
      | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db  273 GGTAAGCAGTTCCTGGCCCCGGCTCAGGCGCCAAAGAACAGATGGAACACGTGAATATGGGCC  214

Qy  179 AACAGGAGATATCTGTGTAAGCAGTTCCTGGCCCC -GCTCAGGCGCCAAAGAACAGATGGTCC  237
      | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db  213 AACAGGAGATATCTGTGTAAGCAGTTCCTGGCCCCGGCTCAGGCGCCAAAGAACAGATGGTCC  154

Qy  238 CCAGATCGGTCCTCGGCCCTCAGCAGTTTCTAGA  270
      | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db  153 CCAGATCGGTCCTCGGCCCTCAGCAGTTTCTAGA  121

RESULT 13
AX166278              AX166278              4148 bp      DNA      linear      PAT 22-JUN-2001
LOCUS                 Sequence 1 from Patent WO0137881.
ACCESSION             AX166278
VERSION               AX166278.1 GI:14546734
KEYWORDS               synthetic construct.
                     synthetic construct
                     artificial sequences.
SOURCE                von Laer,M.D.
REFERENCE              Gene therapy of hiv-positive patients by the expression of
                     membrane-anchored gp41 peptides
                     Patent: WO 0137881-A 1 31-MAY-2001;
                     Heinrich-Pette-Institut (DE)
FEATURES               Location/Qualifiers
   source              1..4148
     /organism="synthetic construct"
     /db_xref="taxon:32630"
     /note="M87 (STHM) - Retroviraaler Vektor MPIN, der 5' von
der IRS-NEO-Kassette ein Insert enthaelt, das fuer
Membran-verankertes T-20-PEPTIDEXodiert"
misc_feature           31..36
     /note="NheI-Schnittstelle"
misc_feature           370..377
     /note="AscI-Schnittstelle"
misc_feature           400..405
     /note="XmaI-Schnittstelle"
misc_feature           696..701
     /note="SacII-Schnittstelle"
misc_feature           1421..1426
     /note="SacII-Schnittstelle"
misc_feature           1429..1436
     /note="NotI-Schnittstelle"
misc_feature           1438..1773
     /note="Bereich im M87 (STHM)-Insert, der fuer
Membran-verankertes T-20-Fusionsprotein einschliesslich
CDS

```

promoter is 993"  
/citation=[3]  
1061. .1076  
/note="multiple cloning site (MCS): contains the unique  
restriction sites XhoI, SacII, NotI"  
1160. .1224  
/note="SV40 small T antigen intron provides splice signal  
for inserts cloned into the MCS"  
1860. .1865  
/note="The polyA signal from the SV40 early region  
provides polyadenylation of inserts cloned in the MCS"  
1881  
1931. .9875  
/note="complete genome of bovine papilloma virus type 1;  
The BPV genome provides for episomal maintenance of the  
vector pBPV when the vector is transfected into mammalian  
cells"  
/citation=[5]  
complement(10647)  
/note="Plasmid origin of replication for replication in E.  
coli. base 10647 represents the first base of the newly  
synthesized strand"  
/direction=left  
complement(11406. .12336)  
/gene="bla"  
complement(11406. .12266)  
/gene="bla"  
/codon\_start=1  
/transl\_table=11  
/product="beta-lactamase"  
/protein\_id="AA57075.1"  
/db\_xref="GI:595689"  
/translation="MSIQHFRVALIPFFAFLPVPFHPETLVKVKDAEDOLGARVGY  
IEDLSKGLDLESPERFPMKSTKVLKGLGAVLSRDVAGQEOIGRIHYSONDLVE  
YSPVTEKHLTDGTVRELCSAALITMSDNTANILITIGSPKELTFLHNGKDVTRL  
DRNEPELNEAIPNDERDTTPAMATTLKLLTGELITLASRQELIDWMDKRVAGPL  
LRSAIPAGWFIADKSGAGERGSGRIIAALGPDPKPSRVIVYITGTSQATMDERNRQIA  
EIGASLIKHW"  
complement(12308. .12336)  
/gene="bla"  
BASE COUNT 3381 a 2852 c 3005 g 3278 t  
ORIGIN

Query Match 94.3%; Score 254.6; DB 12; Length 12516;  
Best Local Similarity 96.7%; Pred. No. 2e-68;  
Matches 260; Conservative 0; Mismatches 9; Indels 0; Gaps 0;  
QY 1 GCTAGCTTAAGTAACGCCATTTTCAGGCATGGGAAATAACATACTGAGATAGAGA 60  
Db 269 GCTAGCTTAAGTAACGCCATTTTCAGGCATGGGAAATAACATACTGAGATAGAGA 210  
61 AGTTCAGATCAAGTTCAGGAACAGAGAAACAGAGATATGGCCAAACAGAGATATCTGT 120  
62 AGTTCAGATCAAGTTCAGGAACAGAGAAACAGAGATATGGCCAAACAGAGATATCTGT 150  
QY 121 GGTAAAGCAGTTCCTGCCCGCTCAGGCCCAGAAAGACAGTGTGGAACAGGAGAAATTTGGGCCAA 180  
Db 149 GGTAAAGCAGTTCCTGCCCGCTCAGGCCCAGAAAGACAGATGAACAGCGTAATTTGGGCCAA 90  
QY 181 ACAGGATATCTGTGTAAGCAGTTCCTGCCCGCTCAGGCCCAGAAACAGAGATGTTCCCA 240  
Db 89 ACAGGATATCTGTGTAAGCAGTTCCTGCCCGCTCAGGCCCAGAAACAGAGATGTTCCCA 30  
QY 241 GATCGGTCGCCCGCTCAGCAGTTCCTAG 269  
Db 29 GATCGGTCGCCCGCTCAGCAGTTCCTAG 1

RESULT 10  
AX449148  
LOCUS  
DEFINITION Sequence 2 from Patent WO0238782.  
ACCESSION AX449148  
AX449148 6575 bp DNA linear PAT 03-JUL-2002

AX449148.1 GI:21697950  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct.  
artificial sequences.  
REFERENCE  
1  
AUTHORS Orberger,G., Hellmuth,K. and Wagener,C.  
TITLE Viral expression system  
JOURNAL Patent: WO 0238782-A 2 16-MAY-2002;  
Armedes Biotechnology GmbH (DE)  
FEATURES  
Location/Qualifiers  
source  
1. .6575  
/organism="synthetic construct"  
/db\_xref="taxon:32630"  
/note="Plasmid pAIM EIB55K, das eine Expressionskassette  
fuer EIB55K enth lt."  
BASE COUNT 1623 a 1566 c 1695 g 1691 t  
ORIGIN

Query Match 91.1%; Score 246; DB 6; Length 6575;  
Best Local Similarity 99.3%; Pred. No. 9.1e-66;  
Matches 268; Conservative 0; Mismatches 0; Indels 2; Gaps 2;  
QY 1 GCTAGCTTAAGTAACGCCATTTTCAGGCATGGGAAATAACATACTGAGATAGAGA 60  
Db 432 GCTAGCTTAAGTAACGCCA-TTTGCAAGCAT-GGAAAAATACATACTGAGATAGAGA 489  
QY 61 AGTTCAGATCAAGTTCAGGAACAGAGAAACAGAGATATGGCCAAACAGAGATATCTGT 120  
Db 490 AGTTCAGATCAAGTTCAGGAACAGAGAAACAGAGATATGGCCAAACAGAGATATCTGT 549  
QY 121 GGTAAAGCAGTTCCTGCCCGCTCAGGCCCAGAAAGACAGTGTGGAACAGGAGAAATTTGGGCCAA 180  
Db 550 GGTAAAGCAGTTCCTGCCCGCTCAGGCCCAGAAAGACAGTGTGGAACAGGAGAAATTTGGGCCAA 609  
QY 181 ACAGGATATCTGTGTAAGCAGTTCCTGCCCGCTCAGGCCCAGAAACAGAGATGTTCCCA 240  
Db 610 ACAGGATATCTGTGTAAGCAGTTCCTGCCCGCTCAGGCCCAGAAACAGAGATGTTCCCA 669  
QY 241 GATCGGTCGCCCGCTCAGCAGTTCCTAG 270  
Db 670 GATCGGTCGCCCGCTCAGCAGTTCCTAG 699

RESULT 11  
MSYMS  
LOCUS MSYMS 2854 bp ss-RNA linear VRL 02-AUG-1993  
DEFINITION Myeloproliferative sarcoma virus proviral v-mos gene, 3' LTR.  
ACCESSION K01683  
VERSION K01683.1 GI:332208  
KEYWORDS c-myc proto-oncogene; coat protein; envelope protein;  
envelope-associated protein; mos oncogene.  
SOURCE Myeloproliferative sarcoma virus proviral DNA.  
ORGANISM Myeloproliferative sarcoma virus  
viruses; Retroviral viruses; Retroviridae; Mammalian type C  
retroviruses; 1-Mammalian type C virus group.  
REFERENCE  
1 (bases 1 to 2854)  
AUTHORS Stacey,A., Abuthnott,C., Kollek,R., Coggin,L. and Ostertag,W.  
TITLE Comparison of myeloproliferative sarcoma virus with Moloney murine  
sarcoma virus variants by nucleotide sequencing and heteroduplex  
analysis  
J. Virol. 50 (3), 725-732 (1984)  
MEDLINE 84216451  
PUBMED 6328002  
COMMENT  
[1] reports an amber mutation (deletion of one 'c') in the envelope  
gene of the MPSV genome, at the junction of the envelope/mos genes.  
What would normally be the twentieth codon is now a termination  
codon. At base 973 an in-phase start codon was found. The MPSV  
sequence is compared with two temperature sensitive mutants derived  
from it, Mo-MuSV variant M1 and Moloney murine leukemia virus  
(Mo-MuLV) by heteroduplex mapping. MPSV wild-type contained a 1 kb  
deletion from the pol gene. It also contained sequences related to  
v-mos genes.

```

|||||
Db 289 GCTAGCTTAAGTAAAGCCATTTTCAAGGCATGGGAAAAATACATACTGAGAAATAGAGA 230
|||||
Qy 61 AGTTACATCAAGTCAAGGAAACAGAGAGAGAGAGATATGGCCAAACAGGATATCTGT 120
|||||
Db 229 AGTTACATCAAGTCAAGGAAACAGAGAGAGAGAGATATGGCCAAACAGGATATCTGT 170
|||||
Qy 121 GGTAAAGCAGTTCTCCCGCTCAGGCGCAAGAGACAGTGTGAACAGGAGAAATGGGCCAA 180
|||||
Db 169 GGTAAAGCAGTTCTCCCGCTCAGGCGCAAGAGACAGTGTGAACAGTGAATTTGGGGCAA 110
|||||
Qy 181 ACAGGATATCTGTTAAGCAGTTCTCCCGCTCAGGCGCAAGAGACAGTGTGGTCCCA 240
|||||
Db 109 ACAGGATATCTGCGTAAAGCAGTTCTCCCGCTCAGGCGCAAGAGACAGTGTGGTCCCA 50
|||||
Qy 241 GATCGGTCCCGCTCAGCAGTCTCTAG 269
|||||
49 GATCGGTCCCGCTCAGCAGTCTCTAG 21

RESULT 8
E28258/c
LOCUS E28258 3263 bp DNA linear PAT 18-JUN-2001
DEFINITION Purification of higher transcription complex from nonhuman
transgenic animal.
ACCESSION E28258
VERSION E28258.1 GI:13025292
KEYWORDS JP 1999004638-A/10.
SOURCE unidentified.
ORGANISM unclassified.
REFERENCE 1 (bases 1 to 3263)
AUTHORS Bernd,K., Eric,B., Michael,M. and Greg,P.
TITLE Purification of higher transcription complex from nonhuman
transgenic animal
JOURNAL Patent: JP 1999004638-A 10 12-JAN-1999;
HOECHST AG
COMMENT OS Unidentified
PN JP 1999004638-A/10
PD 12-JAN-1999
PF 26-MAY-1998 JP 1998144743
PR 26-MAY-1997 DE
PI BERND KIRSCHBAUM,ERIC BERGURUNTO,MICHAEL MEISTERERNST, PI
GREG PORITSU
PC A01A67/027.C12N15/09.C12P21/02//C12P21/08.(C12N15/09,
C12R1.91), PC C12N15/00,
PC (C12N15/00.C12R1.91)
CC Strandedness: Single;
Topology: Linear;
FH Key Location/Qualifiers
FT exon 1..3263.
FEATURES
source
1..3263
/organism="unidentified"
/db_xref="taxon:32644"
BASE COUNT 860 a 794 c 756 g 853 t
ORIGIN
Query Match 94.3%; Score 254.6; DB 6; Length 3263;
Best Local Similarity 96.7%; Pred. No. 1.7e-68;
Matches 260; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

Qy 1 GCTAGCTTAAGTAAAGCCATTTTGAAGGCATGGGAAAAATACATACTGAGAAATAGAGA 60
|||||
Db 289 GCTAGCTTAAGTAAAGCCATTTTGAAGGCATGGGAAAAATACATACTGAGAAATAGAGA 230
|||||
Qy 61 AGTTACATCAAGTCAAGGAAACAGAGAGAGAGATATGGCCAAACAGGATATCTGT 120
|||||
Db 229 AGTTACATCAAGTCAAGGAAACAGAGAGAGAGATATGGCCAAACAGGATATCTGT 170
|||||
Qy 121 GGTAAAGCAGTTCTCCCGCTCAGGCGCAAGAGACAGTGTGAACAGGAGAAATGGGCCAA 180
|||||
Db 169 GGTAAAGCAGTTCTCCCGCTCAGGCGCAAGAGACAGTGTGAACAGTGAATTTGGGGCAA 110
|||||

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Qy 181 ACAGGATATCTGTTAAGCAGTCTCTCCCGCTCAGGCGCAAGAGACAGTGTGGTCCCA 240
|||||
Db 109 ACAGGATATCTGCGTAAAGCAGTCTCTCCCGCTCAGGCGCAAGAGACAGTGTGGTCCCA 50
|||||
Qy 241 GATCGGTCCCGCTCAGCAGTCTCTAG 269
|||||
Db 49 GATCGGTCCCGCTCAGCAGTCTCTAG 21
|||||

RESULT 9
XXU13843/c
LOCUS XXU13843 12516 bp DNA
DEFINITION PBV cloning vector, complete sequence.
ACCESSION U13843
VERSION U13843.1 GI:595688
KEYWORDS bovine papilloma virus; metallothionein I promoter; Moloney sarcoma
virus enhancer; beta-lactamase.
SOURCE unidentified cloning vector.
ORGANISM unidentified cloning vector.
REFERENCE 1 (bases 1 to 12516)
AUTHORS Malone,J.A.
TITLE PBV: An episomally maintained mammalian expression vector based on
bovine papilloma virus
JOURNAL Unpublished (1994)
REFERENCE 2 (bases 1 to 391)
AUTHORS Dhar,R., McClements,W.L., Enquist,L.W. and Vande Woude,G.F.
TITLE Nucleotide sequences of integrated Moloney sarcoma provirus long
terminal repeats and their host and viral junctions
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 77 (7), 3937-3941 (1980)
MEDLINE 81054687
PUBMED 6254003
REFERENCE 3 (bases 692 to 1060)
AUTHORS Glanville,N., Durnam,D.M. and Palmiter,R.D.
TITLE Structure of mouse metallothionein-I gene and its mRNA
JOURNAL Nature 292 (5820), 267-269 (1981)
MEDLINE 81245168
PUBMED 7254320
REFERENCE 4 (bases 939 to 949)
AUTHORS Searle,P.F., Stuart,G.W. and Palmiter,R.D.
TITLE Building a metal-responsive promoter with synthetic regulatory
elements
JOURNAL Mol. and Cell. Biol. 5, 1480-1489 (1985)
REFERENCE 5 (bases 1931 to 9875)
AUTHORS Chen,E.Y., Howley,P.M., Levinson,A.D. and Seeburg,P.H.
TITLE The primary structure and genetic organization of the bovine
papillomavirus type 1 genome
JOURNAL Nature 299 (5883), 529-534 (1982)
MEDLINE 83012974
PUBMED 6289124
REFERENCE 6 (bases 1 to 12516)
AUTHORS Malone,J.A.
TITLE Direct Submission
JOURNAL Submitted (19-AUG-1994) James A. Malone, International Technical
Services, Molecular Biology Reagents Division, Pharmacia Biotech
Inc., 2202 N. Bartlett Ave., Milwaukee, WI 53202-1009, USA

FEATURES
source
1..12516
/organism="unidentified cloning vector"
/db_xref="taxon:45196"
/lab_host="Escherichia coli"
complement(1..391)
/standard_name="Moloney Sarcoma Virus (MSV) enhancer"
/citation=[2]
939..949
/standard_name="metal response element (MRE)"
/citation=[4]
965..970
/feature="mouse metallothionein I"
965..970
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/note="The first base of nascent transcripts from the MT-I

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SOURCE MMSV (Moloney murine sarcoma virus) from transformed mink cells, clone p-600-3.

ORGANISM Moloney murine leukemia virus

REFERENCE Viruses; Retroviral viruses; Retroviridae; Gammaretrovirus.

AUTHORS 1 (bases 1 to 903)

TITLE Dhar,R., McClements,W.L., Enquist,L.W. and Vande Woude,G.F. Nucleotide sequences of integrated Moloney sarcoma provirus long terminal repeats and their host and viral junctions

JOURNAL Proc. Natl. Acad. Sci. U.S.A. 77 (7), 3937-3941 (1980)

MEDLINE 81054687

PUBMED 6254003

COMMENT On Oct 4, 1994 this sequence version replaced gi:331983. [1]: the first and last 11 nucleotides of the terminal repeat sequence (trs) are inverted with respect to each other, and the same four-nucleotide host sequence is found bracketing integrated MMSV. Bases 233-820 represent the 588 bp left trs. See mmsvtrs2 and mmsvtrs3 for the following segments. See also mmsvpro for the complete proviral sequence.

EMBL features not translated to GenBank features:

key	from	to	description
CELL	<1	232	cellular DNA (from mink)
PROVRL	233	>903	proviral DNA
SITE	233	820	terminal repeat
SITE	229	232	repeated again at right end of provirus.

FEATURES

Location/Qualifiers

1..903

/organism="Moloney murine leukemia virus"

/proviral

/db\_xref="taxon:11801"

BASE COUNT 238 a 245 c 215 g 205 t

ORIGIN 352 bases upstream of proviral site.

Query Match 94.7%; Score 255.6; DB 14; Length 903;

Best Local Similarity 96.7%; Pred. No. 7.2e-69;

Matches 261; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

Qy 1 GCTAGCTTAAGTAAGCCATTTCGCAAGGCATGCGGAAATAATACATACTGAGAATAGAGA 60

Db 261 GCTAGCTTAAGTAAGCCATTTCGCAAGGCATGCGGAAATAATACATACTGAGAATAGAGA 320

Qy 61 AGTTTCAGATCAAGTCAGGACAGAACAGAGAGAAATATGGGCCAAACAGGATATCTGT 120

Db 321 AGTTTCAGATCAAGTCAGGACAGAACAGAGAGAAATATGGGCCAAACAGGATATCTGT 380

Qy 121 GGTAAAGCAGTTCTCTGCGCCGCTCAGGGCCCAAGAACAGTGGGAACAGAGAAATTTGGGCCAA 180

Db 381 GGTAAAGCAGTTCTCTGCGCCGCTCAGGGCCCAAGAACAGTGGGAACAGAGAAATTTGGGCCAA 440

Qy 181 ACAGGATATCTGTGTAGCAGTTCTCTGCGCCGCTCAGGGCCCAAGAACAGAGATGGTCCCA 240

Db 441 ACAGGATATCTGTGTAGCAGTTCTCTGCGCCGCTCAGGGCCCAAGAACAGAGATGGTCCCA 500

Qy 241 GATCGGTCCCGCTCAGCAGTTTCTAGA 270

Db 501 GATCGGTCCCGCTCAGCAGTTTCTAGA 530

RESULT 7

LOCUS AX002278/c 3263 bp DNA linear PAT 10-MAR-2000

DEFINITION Sequence 15 from Patent EP0881288.

ACCESSION AX002278

VERSION AX002278.1 GI:7241968

KEYWORDS

SOURCE unidentified.

ORGANISM unidentified.

REFERENCE 1 (bases 1 to 3263)

AUTHORS Berglund,E.D. and Kirschbaum,B.D.

TITLE Purification of higher order transcription complexes from transgenic non-human animals

JOURNAL Patent: EP 0881288-A 15 02-DEC-1998;

FEATURES

Location/Qualifiers

1..3263

/organism="unidentified"

/db\_xref="taxon:32644"

1..3263

BASE COUNT 860 a 794 c 756 g 853 t

ORIGIN

Query Match 94.3%; Score 254.6; DB 6; Length 3263;

Best Local Similarity 96.7%; Pred. No. 1.7e-68;

Matches 260; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

Qy 1 GCTAGCTTAAGTAAGCCATTTCGCAAGGCATGCGGAAATAATACATACTGAGAATAGAGA 60

JOURNAL Proc. Natl. Acad. Sci. U.S.A. 77 (7), 3937-3941 (1980)

MEDLINE 81054687

PUBMED 6254003

FEATURES

Location/Qualifiers

233..903

/organism="Moloney murine sarcoma virus"

/proviral

/db\_xref="taxon:11809"

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/organism="Mustela vison"

/db\_xref="taxon:9667"

229..232

/note="repeated again at right end of provirus"

233..820

/note="terminal repeat"

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Best Local Similarity 96.7%; Pred. No. 7.2e-69;

Matches 261; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

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Db 321 AGTTTCAGATCAAGTCAGGACAGAACAGAGAGAAATATGGGCCAAACAGGATATCTGT 380

Qy 121 GGTAAAGCAGTTCTCTGCGCCGCTCAGGGCCCAAGAACAGTGGGAACAGAGAAATTTGGGCCAA 180

Db 381 GGTAAAGCAGTTCTCTGCGCCGCTCAGGGCCCAAGAACAGTGGGAACAGAGAAATTTGGGCCAA 440

Qy 181 ACAGGATATCTGTGTAGCAGTTCTCTGCGCCGCTCAGGGCCCAAGAACAGAGATGGTCCCA 240

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Qy 241 GATCGGTCCCGCTCAGCAGTTTCTAGA 270

Db 501 GATCGGTCCCGCTCAGCAGTTTCTAGA 530

RESULT 7

LOCUS AX002278/c 3263 bp DNA linear PAT 10-MAR-2000

DEFINITION Sequence 15 from Patent EP0881288.

ACCESSION AX002278

VERSION AX002278.1 GI:7241968

KEYWORDS

SOURCE unidentified.

ORGANISM unidentified.

REFERENCE 1 (bases 1 to 3263)

AUTHORS Berglund,E.D. and Kirschbaum,B.D.

TITLE Purification of higher order transcription complexes from transgenic non-human animals

JOURNAL Patent: EP 0881288-A 15 02-DEC-1998;

FEATURES

Location/Qualifiers

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/organism="unidentified"

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1..3263

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Best Local Similarity 96.7%; Pred. No. 1.7e-68;

Matches 260; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

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QY 61 AGTTCAGATCAAGGTTCAGGAAACAGAGAGAAATATGGGCCAAACAGGATATCTGT 120
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Db 216 ACAGGATATCTGTTAGTACAGTTCCTGCCCGCTCAGGCGCAAGAACAGAGAGAAATGGGCCAA 275
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QY 241 GATCGGTCCCGCCTCAGCAGTTCCTAGA 270
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Db 276 GATCGGTCCCGCCTCAGCAGTTCCTAGA 305
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RESULT 4
LOCUS
DEFINITION
Moloney mouse sarcoma virus right long terminal repeat (as
integrated into mink DNA).
ACCESSION
V01183
VERSION
V01183.1 GI:61641
KEYWORDS
terminal repeat.
SOURCE
Moloney murine sarcoma virus.
ORGANISM
Moloney murine sarcoma virus.
REFERENCE
1 (bases 1 to 646)
Dhar,R., McClements,W.L., Enquist,L.W. and Vande Woude,G.F.
Nucleotide sequences of integrated Moloney sarcoma provirus long
terminal repeats and their host and viral junctions
Proc. Natl. Acad. Sci. U.S.A. 77 (7), 3937-3941 (1980)
JOURNAL
MEDLINE
PUBMED
81054687
FEATURES
Location/Qualifiers
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/focus="proviral DNA integrated into Mink"
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Best Local Similarity 96.7%; Pred. No. 6.9e-69;
Matches 261; Conservative 0; Mismatches 9; Indels 0; Gaps 0;
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RESULT 5
LOCUS
DEFINITION
Moloney murine sarcoma provirus LTR and host sequence, segment 1.
ACCESSION
J02268
VERSION
J02268.1 GI:555246
KEYWORDS
terminal repeat.
SEGMENT
1 of 3
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JOURNAL Patent: WO 0065074-A 1 02-NOV-2000;  
Gesellschaft für Biotechnologische Forschung mbH (GBF) (; DE)  
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LOCUS Plasmid pmBC-2T (expression plasmid-eukaryotic-in vitro) DNA.  
DEFINITION  
ACCESSION X77750  
VERSION X77750.1 GI:456203  
KEYWORDS expression plasmid; Plasmid.  
SOURCE synthetic construct.  
ORGANISM  
artificial sequences.  
REFERENCE  
1 (bases 1 to 3123)  
Dirks, W., Schaper, F. and Hauser, H.  
AUTHORS A new hybrid promoter directs transcription at identical starts  
TITLE points in mammalian cells and in vitro  
JOURNAL Gene 49, 389-390 (1994)  
REFERENCE  
2 (bases 1 to 3123)  
Schaper, F.  
AUTHORS Direct Submission  
TITLE Submitted (14-FEB-1994) F. Schaper, Gesellschaft für  
JOURNAL Biotechnologische Fors., (GBF), Mascheroder Weg 1, 38106  
Braunschweig, FRG  
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QY 61 AGTTTCAGATCAAGTTCAGGAAACAGAGGAGAAATATGGGCCAAACAGGATATCTGT 120  
DB 96 AGTTTCAGATCAAGTTCAGGAAACAGAGGAGAAATATGGGCCAAACAGGATATCTGT 155  
QY 121 GGTAAAGCAGTTCCTGCGCCGCTCAGGCGCCAAAGAACAGTTCGAAACAGGAGAAATTTGGGCCAA 180  
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QY 181 ACAGGATATCTGTGGTAAGCAGTTCCTGCGCCGCTCAGGCGCCAAAGAACAGTTCGAAACAGGAGAAATTTGGGCCAA 240  
DB 216 ACAGGATATCTGTGGTAAGCAGTTCCTGCGCCGCTCAGGCGCCAAAGAACAGTTCGAAACAGGAGAAATTTGGGCCAA 275  
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DB 276 GATCGCGTCCCGCCCTCAGCAGTTTCTAGA 305  
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A70359 8298 bp DNA circular PAT 07-MAY-1999  
LOCUS Sequence 1 from Patent WO9811241.  
DEFINITION  
ACCESSION A70359  
VERSION A70359.1 GI:4774641  
KEYWORDS  
SOURCE Escherichia coli.  
ORGANISM Escherichia coli.  
REFERENCE  
1 (bases 1 to 8298)  
Von, H.I., Bruemmer, W., Burge, C., Riske, E., Dunker, R., Welge, T.,  
AUTHORS Hauser, H. and Mielke, C.  
TITLE OLIGOCISTRONIC EXPRESSION SYSTEM FOR THE PRODUCTION OF HETEROMERIC  
PROTEINS  
JOURNAL Patent: WO 9811241-A 1 19-MAR-1998;  
VON HOEEN IKA (BE)  
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GenCore version 5.1.3  
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- 8: gb\_pl.\*
- 9: gb\_pr.\*
- 10: gb\_ro.\*
- 11: gb\_sts.\*
- 12: gb\_sy.\*
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- 38: em\_sy.\*
- 39: em\_htgo\_hum.\*
- 40: em\_htgo\_mus.\*
- 41: em\_htgo\_other.\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

SUMMARIES

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1	270	100.0	270	6	AX040909 Sequence
2	270	100.0	3123	12	PMBC2TD
3	270	100.0	8298	6	A70359 Sequence 1
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5	255.6	94.7	903	14	MLMTR01
6	255.6	94.7	903	14	REMSV3
7	254.6	94.3	3263	6	AX002278 Sequence
8	254.6	94.3	3263	6	E28258
9	254.6	94.3	12516	12	XX013843
10	246	91.1	6575	6	AX449148
11	223	82.6	2854	14	MSYMO5
12	209	77.4	544	10	S66424
13	208.2	77.1	4148	6	AX166278 Sequence
14	208.2	77.1	9318	6	AR028672
15	206.4	76.4	733	14	MLM3LTR
16	206	76.3	5893	14	REMLV
17	206	76.3	5894	14	AF033812
18	206	76.3	5894	14	MLAPRO
19	202.8	75.1	508	6	AX107878 Sequence
20	202.8	75.1	594	14	MMSLTR1
21	202.8	75.1	635	6	AR016499
22	202.8	75.1	635	6	AR096882
23	202.8	75.1	636	6	E05956
24	202.8	75.1	702	6	I77226
25	202.8	75.1	1200	10	RATDLTR
26	202.8	75.1	1860	10	AX133244S3
27	202.8	75.1	3097	6	AX107931
28	202.8	75.1	4207	6	AX359930 Sequence
29	202.8	75.1	4207	6	AX382144 Sequence
30	202.8	75.1	4210	6	AX359931 Sequence
31	202.8	75.1	4210	6	AX382145 Sequence
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35	202.8	75.1	4661	6	AX359936 Sequence
36	202.8	75.1	4661	6	AX382150
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38	202.8	75.1	4776	6	A95152
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40	202.8	75.1	5109	6	I56770
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42	202.8	75.1	5130	6	AX359935
43	202.8	75.1	5130	6	AX382149
44	202.8	75.1	5176	6	I56771
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ALIGNMENTS

RESULT 1  
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LOCUS AX040909 270 bp DNA linear PAT 23-NOV-2000  
DEFINITION Sequence 1 from Patent WO0065074.  
ACCESSION AX040909  
VERSION AX040909.1 GI:11340531  
KEYWORDS  
SOURCE Myeloproliferative sarcoma virus.  
ORGANISM Myeloproliferative sarcoma virus  
Viruses; Retroviral viruses; Retroviridae; Mammalian type C  
retroviruses; 1-Mammalian type C virus group.  
REFERENCE  
1 (bases 1 to 270)  
AUTHORS Mueller, P., Geserick, C., Schroeder, K. and Hauser, H.  
TITLE Promoter-transactivator system for inducible high-level mammalian  
gene expression with the option of cell growth control